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(54) Title: ORGANOMETALLIC COMPOUNDS AND EMISSION-SHIFTING ORGANIC ELECTROPHORESCENCE

(57) Abstract: Emissive phosphorescent organometallic compounds are described that produce improved electroluminescence, particularly in the blue regions of the visible spectrum. Organic light emitting devices employing such emissive phosphorescent organometallic compounds are also described. Also described is an organic light emitting layer including a host material having a lowest triplet excited state having a decay rate of less than about 1 per second; a guest material dispersed in the host material, the guest material having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 or about 1×10^6 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the guest material.

**ORGANOMETALLIC COMPOUNDS AND EMISSION-SHIFTING ORGANIC
ELECTROPHOSPHORESCENCE**

5

FIELD OF THE INVENTION

The present invention is directed to phosphorescence based organic light emitting devices that have improved electroluminescent characteristics.

10

BACKGROUND OF THE INVENTION

The technology of organic light emitting diodes (OLEDs) is undergoing rapid development. OLEDs originally utilized the electroluminescence produced from 15 electrically excited molecules that emitted light from their singlet states. Such radiative emission from a singlet excited state is referred to as fluorescence. More recent work has demonstrated that higher power efficiency OLEDs can be made using molecules that emit light from their triplet state, defined as phosphorescence.

20

Such electrophosphorescence makes it possible for phosphorescent OLEDs to have substantially higher quantum efficiencies than are possible for OLEDs that only produce fluorescence. This is based on the understanding that the excitons created in an OLED are produced, according to simple statistical arguments as well as experimental measurements, approximately 75% as triplet excitons and 25% as singlet excitons. The

triplet excitons more readily transfer their energy to triplet excited states that can produce phosphorescence whereas the singlet excitons typically transfer their energy to singlet excited states that can produce fluorescence. Since the lowest emissive singlet excited state of an organic molecule is typically at a slightly higher energy than the lowest triplet excited state, the singlet excited state may relax, by an intersystem crossing process, to the emissive triplet excited state. This means that all the exciton excitation energy may be converted into triplet state excitation energy, which then becomes available as phosphorescent emission. Thus, electrophosphorescent OLEDs have a theoretical quantum efficiency of 100%, since all the exciton excitation energy can become available as electrophosphorescence.

As a consequence, since the discovery that phosphorescent materials could be used in an OLED, *Baldo et al., "Highly Efficient Phosphorescent Emission from Organic Electroluminescent Devices"*, *Nature*, vol. 395, 151-154, 1998, there is now much interest in finding more efficient electrophosphorescent materials.

Typically phosphorescent emission from organic molecules is less common than fluorescent emission. However, phosphorescence can be observed from organic molecules under an appropriate set of conditions. Organic molecules coordinated to lanthanide elements often phosphoresce from excited states localized on the lanthanide metal. The europium diketonate complexes illustrate one group of these types of species. Organic phosphorescence is also often observed in molecules containing heteroatoms with unshared pairs of electrons at very low temperatures. Benzophenone and 2,2'-bipyridine

are such molecules. Phosphorescence can be enhanced over fluorescence by confining, preferably through bonding, the organic molecule in close proximity to an atom of high atomic number. This phenomenon, called the heavy atom effect, is created by a mechanism known as spin-orbit coupling. A related phosphorescent transition is a metal-
5 to-ligand charge transfer (MLCT) that is observed in molecules such as tris(2-phenylpyridine)iridium(III).

However, molecules that phosphoresce from MLCT states typically emit light that is of lower energy than that observed from the unbound organic ligand. This
10 lowering of emission energy makes it difficult to develop organic molecules that phosphoresce in the technologically useful blue and green colors of the visible spectrum where the unperturbed phosphorescence typically occurs.

It would be desirable if more efficient electrophosphorescent materials
15 could be found, particularly materials that produce their emission in the blue region of the spectrum.

The realization of highly efficient blue, green and red electrophosphorescence is a requirement for portable full color displays and white lighting
20 applications with low power consumption. Recently, high-efficiency green and red organic electrophosphorescent devices have been demonstrated which harvest both singlet and triplet excitons, leading to internal quantum efficiencies (η_{int}) approaching 100%. See Baldo, M. A., O'Brien, D. F., You, Y., Shoustikov, A., Sibley, S., Thompson, M. E., and

Forrest, S. R., *Nature (London)*, **395**, 151-154 (1998); Baldo, M. A., Lamansky, S., Burrows, P. E., Thompson, M. E., and Forrest, S. R., *Appl. Phys. Lett.*, **75**, 4-6 (1999); Adachi, C., Baldo, M. A., and Forrest, S. R., *App. Phys. Lett.*, **77**, 904-906, (2000); Adachi, C., Lamansky, S., Baldo, M. A., Kwong, R. C., Thompson, M. E., and Forrest, S. 5 R., *App. Phys. Lett.*, **78**, 1622-1624 (2001); and Adachi, C., Baldo, M. A., Thompson, M. E., and Forrest, S. R., *Bull. Am. Phys. Soc.*, **46**, 863 (2001). Using a green phosphor, *fac*-tris(2-phenylpyridine)iridium ($\text{Ir}(\text{ppy})_3$), in particular, an external quantum efficiency (η_{ext}) of $(17.6 \pm 0.5)\%$ corresponding to an internal quantum efficiency of $>85\%$, was realized using a wide energy gap host material, 3-phenyl-4-(1'-naphthyl)-5-phenyl-1,2,4-triazole (TAZ). See Adachi, C., Baldo, M. A., Thompson, M. E., and Forrest, S. R., *Bull. Am. Phys. Soc.*, **46**, 863 (2001). Most recently, high- efficiency ($\eta_{\text{ext}} = (7.0 \pm 0.5)\%$) red electrophosphorescence was demonstrated employing bis(2-(2'-benzo[4,5-a]thienyl)pyridinato-N, C³) iridium (acetylacetone) [$\text{Btp}_2\text{Ir}(\text{acac})$]. See Adachi, C., Lamansky, S., Baldo, M. A., Kwong, R. C., Thompson, M. E., and Forrest, S. R., *App. 10 Phys. Lett.*, **78**, 1622-1624 (2001).

In each of these latter cases, high efficiencies are obtained by energy transfer from both the host singlet and triplet states to the phosphor triplet, or via direct trapping of charge on the phosphor, thereby harvesting up to 100% of the excited states. 15 This is a significant improvement over what can be expected using fluorescence in either small molecule or polymer organic light emitting devices (OLEDs). See Baldo, M. A., O'Brien, D. F., Thompson, M. E., and Forrest, S. R., *Phys. Rev. B* **60**, 14422-14428 (1999); Friend, R. H., Gymer, R. W., Holmes, A. B., Burroughes, J. H., Marks, R. N.,

Taliani, C., Bradley, D. D. C., Dos Santos, D. A., Bredas, J. L., Logdlund, M., Salaneck, W. R., *Nature (London)*, **397**, 121-128 (1999); and Cao, Y., Parker, I. D., Yu, G., Zhang, C., and Heeger, A. J., *Nature (London)*, **397**, 414-417 (1999). In either case, these transfers entail a resonant, exothermic process. As the triplet energy of the phosphor increases, it becomes less likely to find an appropriate host with a suitably high energy triplet state. See Baldo, M. A., and Forrest, S. R., *Phys. Rev. B* **62**, 10958-10966 (2000). The very large excitonic energies required of the host also suggest that this material layer may not have appropriate energy level alignments with other materials used in an OLED structure, hence resulting in a further reduction in efficiency. To eliminate this competition between the conductive and energy transfer properties of the host, a route to efficient blue electrophosphorescence may involve the endothermic energy transfer from a near resonant excited state of the host to the higher triplet energy of the phosphor. See Baldo, M. A., and Forrest, S. R., *Phys. Rev. B* **62**, 10958-10966 (2000); Ford, W. E., Rodgers, M. A. J., *J. Phys. Chem.*, **96**, 2917-2920 (1992); and Harriman, A.; Hissler, M.; Khatyr, A.; Ziessel, R. *Chem. Commun.*, 735-736 (1999). Provided that the energy required in the transfer is not significantly greater than the thermal energy, this process can be very efficient.

Organic light emitting devices (OLEDs), which make use of thin film materials that emit light when excited by electric current, are expected to become an increasingly popular form of flat panel display technology. This is because OLEDs have a wide variety of potential applications, including cellphones, personal digital assistants (PDAs), computer displays, informational displays in vehicles, television monitors, as well

as light sources for general illumination. Due to their bright colors, wide viewing angle, compatibility with full motion video, broad temperature ranges, thin and conformable form factor, low power requirements and the potential for low cost manufacturing processes, OLEDs are seen as a future replacement technology for cathode ray tubes (CRTs) and liquid crystal displays (LCDs), which currently dominate the growing \$40 billion annual electronic display market. Due to their high luminous efficiencies, electrophosphorescent OLEDs are seen as having the potential to replace incandescent, and perhaps even fluorescent, lamps for certain types of applications.

10 Light emission from OLEDs is typically via fluorescence or phosphorescence. As used herein, the term "phosphorescence" refers to emission from a triplet excited state of an organic molecule and the term fluorescence refers to emission from a singlet excited state of an organic molecule.

15 Successful utilization of phosphorescence holds enormous promise for organic electroluminescent devices. For example, an advantage of phosphorescence is that all excitons (formed by the recombination of holes and electrons in an EL), which are formed either as a singlet or triplet excited state, may participate in luminescence. This is because the lowest singlet excited state of an organic molecule is typically at a slightly higher energy than the lowest triplet excited state. This means that, for typical phosphorescent organometallic compounds, the lowest singlet excited state may rapidly decay to the lowest triplet excited state from which the phosphorescence is produced. In contrast, only a small percentage (about 25%) of excitons in fluorescent devices are

capable of producing the fluorescent luminescence that is obtained from a singlet excited state. The remaining excitons in a fluorescent device, which are produced in the lowest triplet excited state of an organic molecule, are typically not capable of being converted into the energetically unfavorable higher singlet excited states from which the fluorescence is produced. This energy, thus, becomes lost to radiationless decay processes that heat-up
5 the device.

SUMMARY OF THE INVENTION

10 The present invention is directed to emissive phosphorescent organometallic compounds that produce improved electroluminescence, organic light emitting devices employing such emissive phosphorescent organometallic compounds, and methods of fabricating such organic light emitting devices.

15 Specific embodiments of the present invention are directed to OLEDs using emissive phosphorescent organometallic compounds that produce improved electrophosphorescence in the blue region of the visible spectrum.

The present invention is directed, in addition, to a method of selecting
20 organometallic compounds that have improved electroluminescent properties, for example, in the blue region of the visible spectrum.

The present invention is also directed to an organic light emitting layer

including a host material and a guest material dispersed in the host material, the guest material having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 or about 1×10^6 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the guest material. The sum of the radiative and non-radiative decay rates of the host triplet is preferably not greater than about $5 \times 10^3/\text{sec}$, and more preferably, not greater than about $1 \times 10^3/\text{sec}$.

The present invention is also directed to an organic light emitting layer including a host material having a lowest triplet excited state having a decay rate of less than about 1 per second; a guest material dispersed in the host material, the guest material having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 or about 1×10^6 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the guest material.

BRIEF DESCRIPTION OF THE DRAWINGS

For the purpose of further illustrating the invention, representative embodiments are shown in the accompanying figures, it being understood that the invention is not intended to be limited to the precise arrangements and instrumentalities shown.

Figures 9a through 9g show the chemical structures of representative organometallic compounds described herein.

Figure 10 shows the emission spectrum of both Pt(ppy)₂ and Pt(ppy)₂Br₂.

5 The former gives green emission, partly from MLCT transitions, and the latter gives blue emission, predominantly from a triplet π - π * transition. The structure observed for the Pt(ppy)₂Br₂ spectrum is consistent with ligand-centered emission. The luminescent lifetimes for the two complexes are 4 and 150 microseconds.

10 Figure 11 is a plot showing the emission spectra of (ppy)AuCl₂ and (ppy)Au(2,2'biphenylene). Both emit from ligand triplet π - π * transitions.

Figure 12 is a CIE diagram providing the coordinates of (C-N)Pt(acac) complexes. All coordinates are based on solution photoluminescent measurements except 15 for 4,5-F₂ppy-EL, which corresponds to the electroluminescent spectrum. The Ir(ppy)₃ is an electroluminescent spectrum as well.

Figure 13 is a plot depicting the photoluminescent emission spectra of (4,6-F₂ppy)Pt(acac) at room temperature (RT) and at 77 K. Also shown are the excitation 20 spectra taken at 77 K and the absorbance spectra taken at room temperature for the same complex.

Figure 14 illustrates the normalized photoluminescent emission spectra of

(ppy)Pt(acac), (4,5 dfppy)Pt(acac), and (4,5 dfppy)Pt(pico).

Figure 15 illustrates the normalized photoluminescent emission spectra of typPy(acac), bzqPt(acac), and btpPt(acac).

5

Figure 16 illustrates the normalized electroluminescent emission spectra for OLEDs prepared with (2-(4,5-F₂phenyl)pyridinato)platinum(acetyl acetonate). The OLEDs had a ITO/PVK-PBD-dopant/Alq₃/Mg-Ag layer structure. The PVK layer was deposited as a single, homogeneous layer by spin coating. PVK = polyvinylcarbaazole and PBD = (4-biphenyl)(4-tertbutyl)oxidiazole. The Alq₃ and Mg-Ag layers were deposited by thermal evaporation. The OLED had an external efficiency of 1.3 % and a turn on voltage of 5 Volts. The spectra of the EL output as well as the PL signal are shown.

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Figure 17 shows the molecular structures of some materials studied for the present invention and a view of the triplet dynamics in a guest-host system of the present invention.

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Figure 18 shows the structure of the electroluminescent devices used to observe the transient response of triplet diffusion in organic host materials.

Figure 19 shows the phosphorescent spectra of TPD, BCP, CBP and Ir(ppy)₃ with PtOEP according to the present invention.

Figure 20 shows the transient response of four phosphorescent guest-host systems according to the present invention.

Figure 21 shows the electroluminescent response of 8% Ir(ppy)₃ in TPD.

5

Figure 22 shows the normalized phosphorescent transients for PtOEP in Alq₃ recorded at 650 nm for diffusion distances of (a) 200 Å, (b) 400 Å, (c) 600 Å and (d) 800 Å.

10 Figure 23 shows the exciton current incident in the phosphorescent zone for diffusion distances (a) 200 Å, (b) 400 Å, (c) 600 Å and (d) 800 Å.

DETAILED DESCRIPTION

15 The present invention will now be described in detail for specific preferred embodiments of the invention. These embodiments are intended only as illustrative examples and the invention is not to be limited thereto.

20 The phosphorescent organometallic compounds of the present invention are comprised of: (a) a heavy transition metal such as Ir, but not limited to Ir, which produces efficient phosphorescent emission at room temperature from a mixture of MLCT and $\pi - \pi^*$ ligand states; (b) wherein the metal is bound to at least one mono-anionic, bidentate, carbon-coordination ligand substituted with electron donating and/or electron withdrawing

substituents that shift the emission, relative to the un-substituted ligand, to either the blue, green or red region of the visible spectrum; and (c) wherein the metal is bound to at least one non-mono-anionic, bidentate, carbon-coordination ligand, which may be substituted or un-substituted, that causes the emission to have a well defined vibronic structure.

5

A carbon-coordination ligand is a ligand that is bound to the metal atom via a carbon-metal bond. In view of what one skilled in the art might view as a strict definition of organometallic compounds, such as described in *Inorganic Chemistry*, by Gary L. Miessler and Donald A. Tarr, 2nd edition, Prentice Hall, 1999, the compounds of the present invention are referred to herein as organometallic compounds since these compounds include a metal-carbon bond.

The phosphorescent organometallic compounds of the present invention have at least one carbon-coordination ligand wherein the at least one carbon-coordination ligand is a mono-anionic ligand. That is, the metal atom is bound to only one carbon atom of the at least one carbon-coordination ligand. Furthermore, the at least one mono-anionic, carbon-coordination ligand of the present invention is a bidentate ligand. A bidentate ligand has two points at which it attaches to a central atom, in this case, the metal atom. Thus, the phosphorescent organometallic compounds of the present invention have at least one mono-anionic, bidentate, carbon-coordination ligand.

The at least one mono-anionic, bidentate, carbon-coordination ligand of the present invention is substituted with electron donating and/or electron withdrawing

substituents that shift the emission, relative to the un-substituted ligand, to either the blue, green or red region of the visible spectrum. The particular substituents used on particular ligands will depend upon the desired shift in emission. Generic representative examples of the at least one mono-anionic, bidentate, carbon-coordination ligand of the present invention are listed in Figures 5a, 5b and 5c. In addition, two specific examples of the at least one mono-anionic, bidentate, carbon-coordination ligand of the present invention with specific substituents are listed in Figure 5d. As can be seen in Figures 5a, 5b and 5c, the at least one mono-anionic, bidentate, carbon-coordination ligand of the present invention can form a cyclometallated ring that includes the organometallic carbon-metal bond and a dative bond between the metal atom and a nitrogen, sulfur or oxygen group.

The carbon atom that is bound to the metal may be present as part of a substituted or unsubstituted, saturated hydrocarbon; a substituted or unsubstituted, aromatic system, for example, phenylene or naphthalene compounds; or a substituted or unsubstituted heterocyclic system, which might include, for example, substituted or unsubstituted thiophenes, furans, pyridines and pyrroles. The group in the cyclometallated ring that forms a dative bond with the metal atom may be independently selected also to include a substituted or unsubstituted, saturated hydrocarbon; a substituted or unsubstituted, aromatic system, for example, phenylene or naphthalene compounds; or a substituted or unsubstituted heterocyclic system, which might include, for example, thiophenes, furans, pyridines and pyrroles. One of these aforementioned groups must be substituted, however, because the at least one mono-anionic, bidentate, carbon-coordination ligand of the present invention is substituted with electron donating and/or electron withdrawing substituents that shift the emission relative to the un-substituted ligand.

The preferred metals of the present invention are metals that can provide strong spin-orbit coupling of the metal atom with the at least one mono-anionic, bidentate, carbon-coordination ligand. Such metals include, in particular, the heavy metals having an atomic number of at least 72, such as Os, Ir, Pt and Au, with Ir and Pt being particularly preferred metals.

5

In addition to being bound to at least one mono-anionic, bidentate, carbon-coordination ligand, the metal atom of the organometallic compounds of the present invention is also bound to at least one non-mono-anionic, bidentate, carbon-coordination ligand. The at least one non-mono-anionic, bidentate, carbon-coordination ligand of the present invention is either not mono-anionic, not bidentate, not a carbon-coordination ligand, or some combination thereof. The at least one non-mono-anionic, bidentate, carbon-coordination ligand causes the emission to have a well defined vibronic structure, and generic representative examples thereof are listed in Figures 6a and 6b. In addition, specific examples of the at least one non-mono-anionic, bidentate, carbon-coordination ligand of the present invention with specific substituents are listed in Figure 6c.

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In one embodiment of the organometallic compounds of the present invention, the organometallic compound includes, in particular, a metal atom bound to a single carbon-coordination ligand, wherein the carbon-coordination ligand is a mono-anionic carbon-coordination ligand. In particular, the metal atom is bound to only one carbon atom of the carbon-coordination ligand. Thus, while the organometallic compounds that are used in the OLEDs of the present invention include more than one

ligand, in this embodiment of the present invention only one ligand is a carbon-coordination ligand. Thus, in this embodiment of the present invention the organometallic compounds include only one carbon-metal bond.

5 In this same embodiment of the present invention, the carbon-coordination ligand is preferably selected from those ligands that exhibit strong charge transfer absorption characteristics, for example, a molar absorptivity of at least 1,000 L/mole-cm, preferably, at least about 2,000-4,000 L/mole-cm. Such absorption bands involve transfer of electrons from molecular orbitals that are primarily ligand in character to orbitals that
10 are primarily metal in character or, alternatively, from orbitals that are primarily metal in character to molecular orbitals that are primarily ligand in character. Miessler and Tarr.
Such an excitation mechanism results in a charge transfer transition that may be
designated as a ligand-to-metal charge transfer (LMCT) or as a metal-to-ligand charge
transfer (MLCT), respectively. The former may be characterized as a partial reduction of
15 the metal atom and the latter as a partial oxidation of the metal atom.

Selection of a carbon-coordination ligand to give a high molar absorptivity of the organometallic compound results in an organometallic compound that is capable of providing highly efficient electroluminescence when used in an OLED. However, rather
20 than functioning as strongly absorbing species in the OLED, such organometallic compounds have highly emissive excited states that are produced when a voltage is applied across the OLED. The high molar absorptivities of such ligands may be used to select ligands that produce highly efficient electroluminescence in an OLED. Such

ligands may be selected to have empty pi-symmetry orbitals on the ligands that become acceptor orbitals upon absorption of light.

In this same embodiment of the present invention, the ligand is preferably selected, in particular, so as to give a strong metal-to-ligand charge transfer (MLCT) absorption band. Such ligands are selected to have empty anti-bonding π^* orbitals on the ligands that become acceptor orbitals upon absorption of light. As representative embodiments of the present invention, the carbon-coordination ligand may be selected from the class of materials such as described, for example, in Comprehensive Coordination Chemistry, Vols. 1-7, G. Wilkinson, Ed., Pergamon Press, 1987.

In this same embodiment of the present invention, in addition to being bound to a single mono-anionic carbon-coordination ligand, the metal atom of the organometallic compound is also bound to one or more additional ligands, each of which are all non-carbon-coordination ligands. A non-carbon-coordination ligand is one that does not form any metal-carbon bonds with the metal atom of the organometallic compound. Preferably, in this same embodiment of the present invention, a metal to ligand charge transfer complex (MLCT) is employed, where the non-carbon-coordination ligands are preferably ligands having a strong electrophilic character such that the ligands draw electrons away from the metal atom. Representative non-carbon-coordination ligands may also be selected, for example, from Comprehensive Coordination Chemistry, Vols. 1-7, G. Wilkinson, Ed., Pergamon Press, 1987.

Without intending to be limited to the theory of how the present invention works, it is believed that the improved electroluminescent properties that are observed for the OLEDs of the present invention may be attributed to a combination of factors. For example, it is believed that selection of heavy metals that are capable of forming metal-to-ligand charge transfer (MLCT) states with carbon-coordination ligands that have empty π^* orbitals, such phosphorescent materials produce highly efficient electrophosphorescent OLEDs. The electroluminescence from representative organometallic compounds of the present invention shows a vibronic fine structure that indicates that the emission is from an excited state that has a wave function represented by a mixture of the MLCT state of the 5 organometallic compound and the excited triplet state of the carbon-coordination ligand. Since the radiative emission is from a triplet excited state, the emission is referred to as phosphorescence.

It is further believed a higher energy radiative emission may be achieved by 15 including electron-withdrawing groups on the carbon-coordination ligand and/or by selecting the non-carbon-coordination ligand to have a strong electron withdrawing character. Without being limited to the precise theory of how the higher energy radiative emissive may be achieved, it is believed that the electron-withdrawing groups tend to remove electron density from the highest occupied molecular orbitals (HOMO) that 20 include the ligand and the metal atom, thus altering the relative energy levels of the ground state and the excited state such that the overall MLCT transition energy from the ground state to the excited state increases. The preferred organometallic compounds for certain embodiments of the present invention include, thus, strong electron-withdrawing groups

on the carbon-coordination ligand and/or non-carbon-coordination ligands having a strong electron-withdrawing character.

Another aspect of the present invention relates to the discovery that OLEDs incorporating emissive organometallic compounds having a single mono-anionic carbon-coordination ligand have substantially higher external quantum efficiencies than compounds with bis-substituted carbon-coordination ligands. For example, the compound having the chemical structure [(ppy)Pt(acac)] was found to produce strong photophosphorescence at room temperature. In contrast, a compound having the structure [Pt(ppy)₂] was found not to produce any visible photophosphoresce at room temperature.

The phosphorescent organometallic compounds of the present invention have at least one mono-anionic, bidentate, carbon-coordination ligand and at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to a heavy transition metal, such as Ir. For example, representative embodiments of the phosphorescent organometallic compounds of the present invention may be illustrated by combining at least one of the mono-anionic, bidentate, carbon-coordination ligands of Figures 5a, 5b, 5c and 5d, at least one of the non-mono-anionic, bidentate, carbon-coordination ligands of Figures 6a, 6b and 6c, and a heavy transition metal, such as Ir. Representative examples of the phosphorescent organometallic compounds of the present invention, along with their emission spectra, are shown in Figures 7a – 7r, it being understood that the present invention is not intended to be limited to the representative examples shown.

The preparation of these representative examples of the phosphorescent organometallic compounds of the present invention shown in Figures 7a – 7r was accomplished as follows.

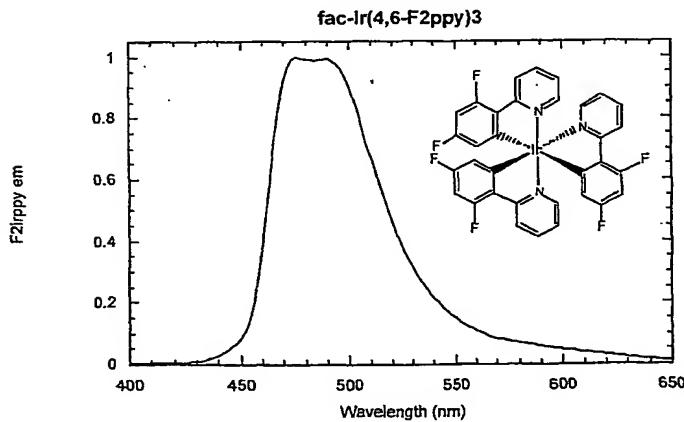
Synthesis of 2-(4,6-difluorophenyl)pyridine

The 2-(4,6-difluorophenyl)pyridine ligand precursor was prepared by Suzuki coupling of 4,6-difluorophenylboronic acid (Frontier Chemical) with 2-bromopyridine (Aldrich) in 1,2-dimethoxyethane using a $\text{Pd}(\text{OAc})_2/\text{PPh}_3$ catalyst and K_2CO_3 base as per.

5 *Synlett, 1999, 1, 45–48.*

Synthesis of fac-tris(2-(4,6-difluorophenyl)pyridinato-N,C^{2'}) Iridium (III).

The $\text{Ir}(\text{acac})_3$ complex was treated with 6 eq of 2-(4,6-difluorophenyl)pyridine in glycerol at 180°C under inert gas atmosphere for 16 hours. After cooling down to room temperature, water was added to the reaction mixture in order to precipitate the crude product. the solvent was removed under reduced pressure and the crude yellow product was washed with methanol to remove any unreacted picolinic acid. The crude product was flash chromatographed using a silica:dichloromethane column to yield ca. 75% of the pure yellow fac-tris(2-(4,6-difluorophenyl)pyridinato-N,C^{2'}) iridium (III) after solvent evaporation and drying.



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Synthesis of [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂.

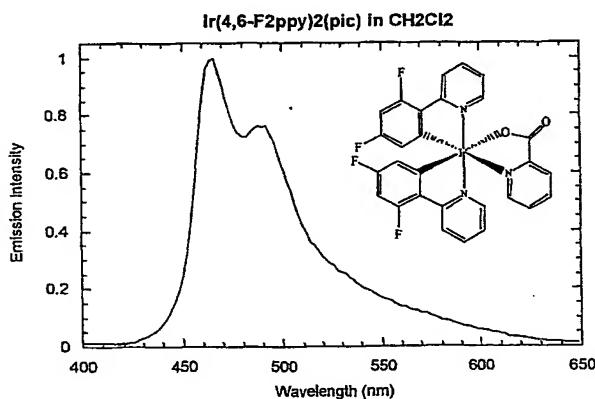
All procedures involving $\text{IrCl}_3 \cdot \text{H}_2\text{O}$ or any other Ir(III) species were carried out in inert gas atmosphere in spite of the air stability of the compounds, the main concern being their oxidative stability and stability of intermediate complexes at high temperatures used in the reactions. The cyclometalated Ir(III) μ -chloro bridged dimer of a general formula

20

C-N₂Ir(μ-Cl)₂IrC-N₂ was synthesized by heating a mixture of IrCl₃•nH₂O with 4 eq. of 2-(4,6-difluorophenyl)pyridine in 2-ethoxyethanol at 130°C for 16 hr. The product was isolated by addition of water followed by filtration and methanol wash. Yield 90%.

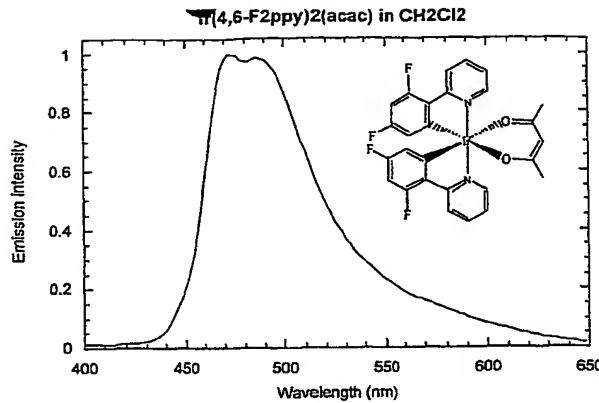
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(picolinate).

5 The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex was treated with 2 eq of picolinic acid in refluxing 1,2-dichloroethane under inert gas atmosphere for 16 hours. After cooling down to room temperature, the solvent was removed under reduced pressure and the crude yellow product was washed with methanol to remove any unreacted picolinic acid. The crude product was flash chromatographed using a
10 silica:dichloromethane column to yield ca. 75% of the pure yellow (C-N₂)₂Ir(pic) after solvent evaporation and drying.



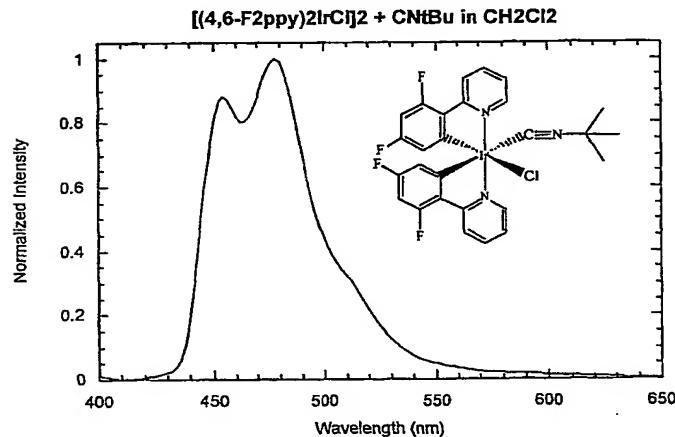
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(acetyl acetonate).

15 The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex was treated with 5 eq of 2,4-pentadione and 10 eq of Na₂CO₃ in refluxing 1,2-dichloroethane under inert gas atmosphere for 16 hours. After cooling down to room temperature, the solvent was removed under reduced pressure and the crude yellow product was washed with methanol. The crude product was flash chromatographed using a silica:dichloromethane column to yield ca. 75% of the pure yellow (C-N₂)₂Ir(acac) after solvent evaporation and drying.



Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C²')(tert-butylisocyanide) chloride.

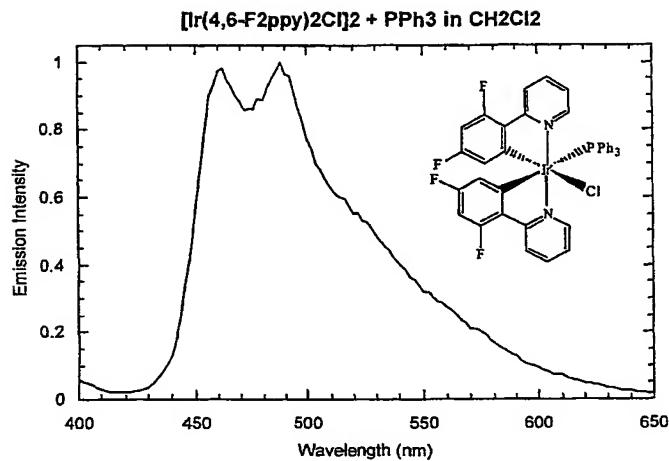
The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with



5 an excess of *tert*-butylisocyanide in 2 mL CH₂Cl₂ solution for 16 hours.

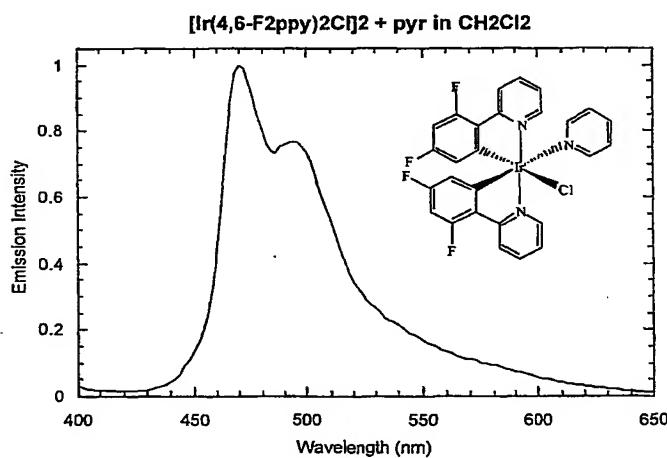
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C²')(triphenylphosphine) chloride.

10 The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with an excess of triphenylphosphine in 2 mL CH₂Cl₂ solution for 16 hours.



Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(pyridine) chloride.

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with

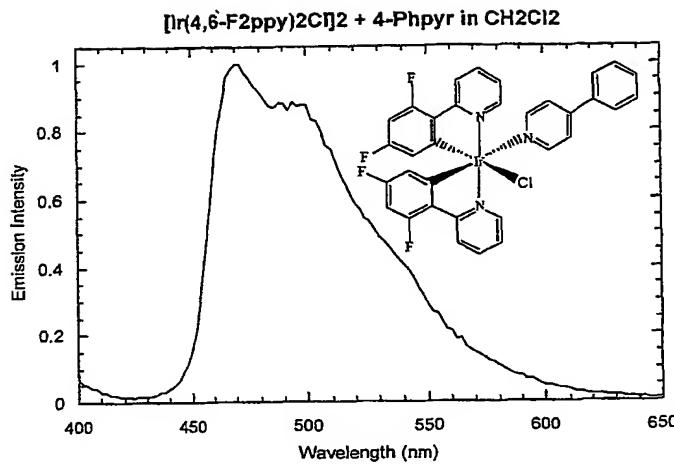


an excess of pyridine in 2 mL CH₂Cl₂ solution for 16 hours.

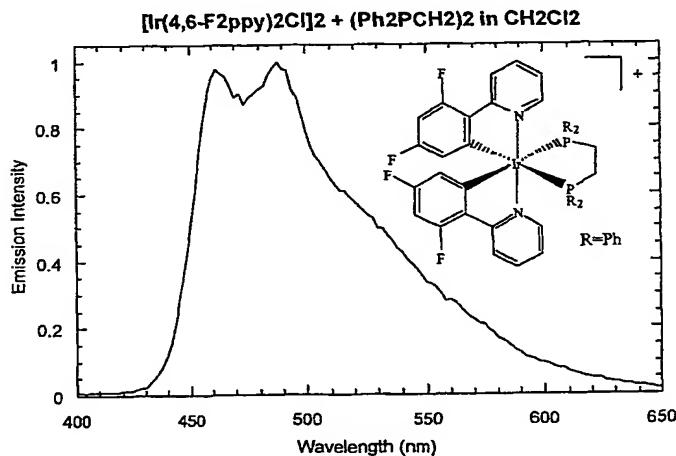
5

Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(4-phenylpyridine) chloride.

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with an excess of 4-phenylpyridine in 2 mL CH₂Cl₂ solution for 16 hours.



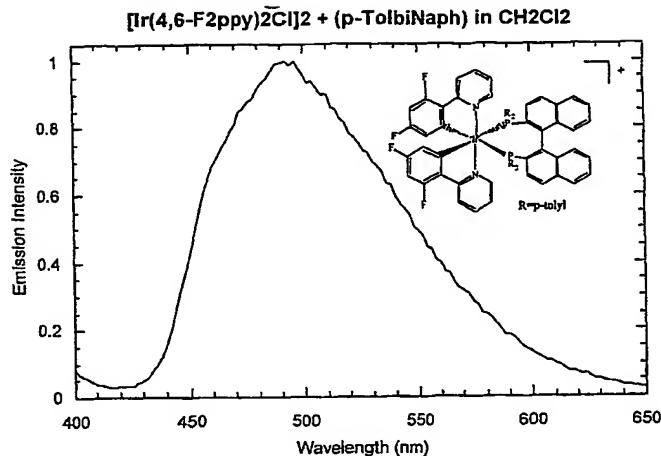
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(1,2-bis(diphenylphosphino) ethane) chloride.



The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with
5 an excess of 1,2-bis(diphenylphosphino)ethane in 2 mL CH₂Cl₂ solution for 16 hours.

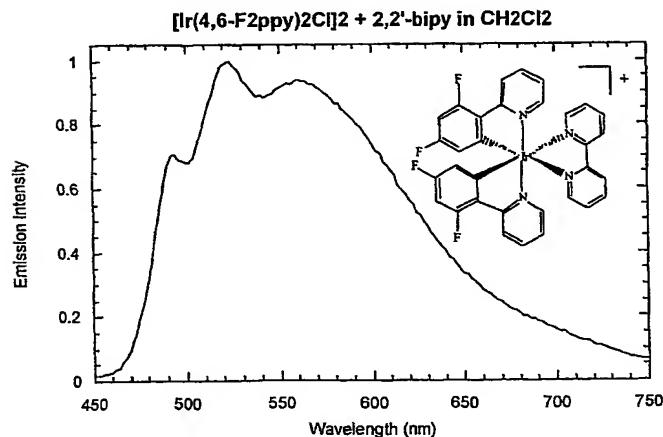
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(*(R*)-(+)2,2'-bis(di-p-tolylphosphino)-1,1'-binaphthyl) chloride.

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with
10 an excess of (*R*)-(+)2,2'-bis(di-p-tolylphosphino)-1,1'-binaphthyl in 2 mL CH₂Cl₂ solution for 16 hours.



Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(2,2'-bipyridine) chloride.

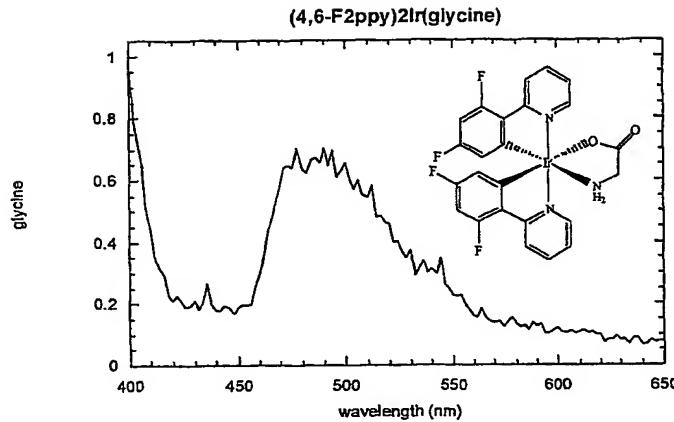
The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂ complex (ca. 0.002 g) was treated with



an excess of 2,2'-bipyridine in 2 mL CH₂Cl₂ solution for 16 hours.

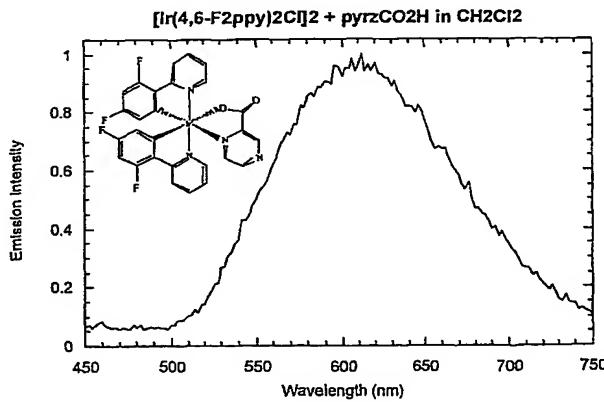
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(glycine).

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with an excess of glycine in 2 mL CH₂Cl₂ solution for 16 hours.



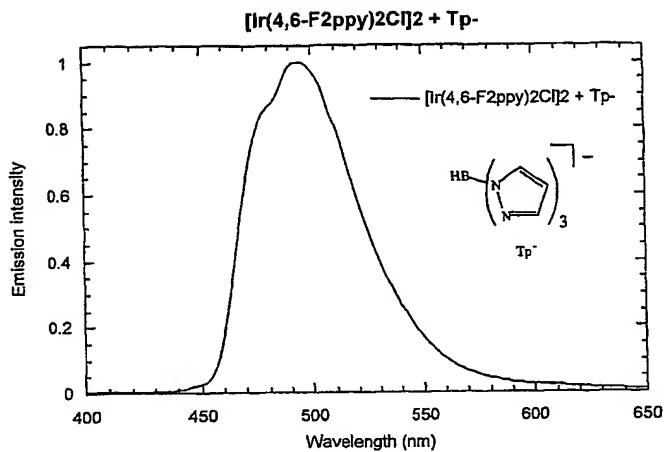
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(pyrazinecarboxylate).

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂ complex (ca. 0.002 g) was treated with
5 an excess of pyrazinecarboxylic acid in 2 mL CH₂Cl₂ solution for 16 hours.



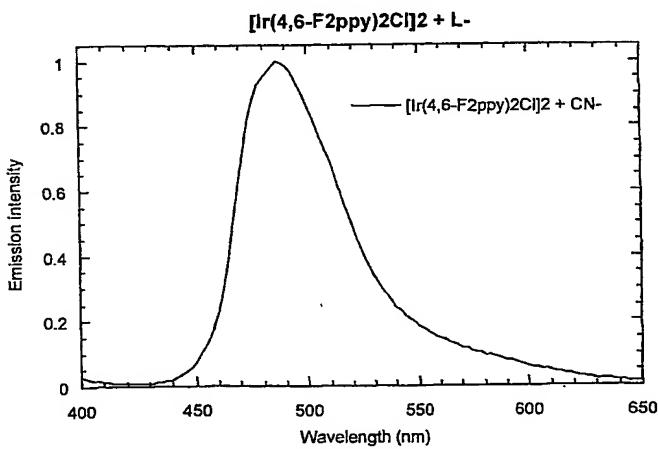
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(tris(pyrazolyl)borate).

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂ complex (ca. 0.002 g) was treated with
10 an excess of potassium tris(pyrazolyl)borate in 2 mL CH₂Cl₂ solution for 16 hours.



Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C²⁻)(cyanide).

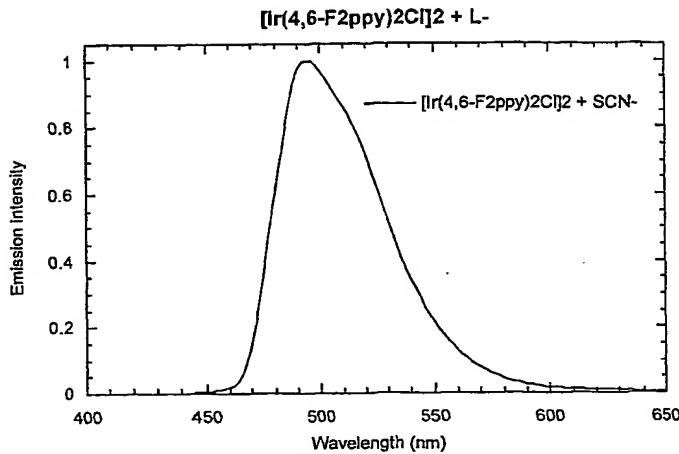
The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with an excess of potassium cyanide in 2 mL acetone solution for 16 hours.



5

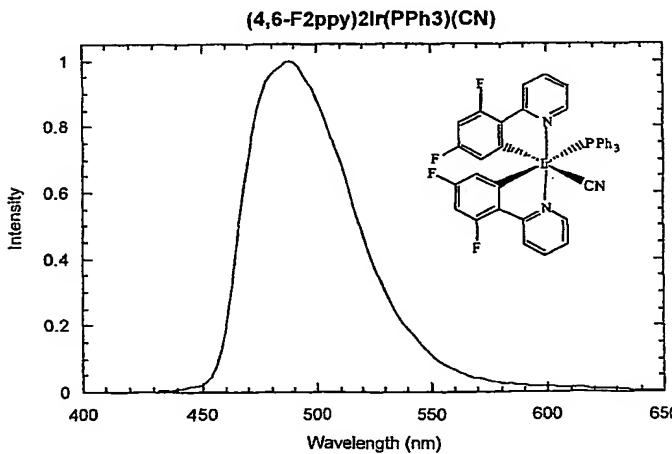
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C²⁻)(thiocyanide).

The [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂complex (ca. 0.002 g) was treated with an excess of sodium thiocyanide in 2 mL CH₂Cl₂ solution for 16 hours.



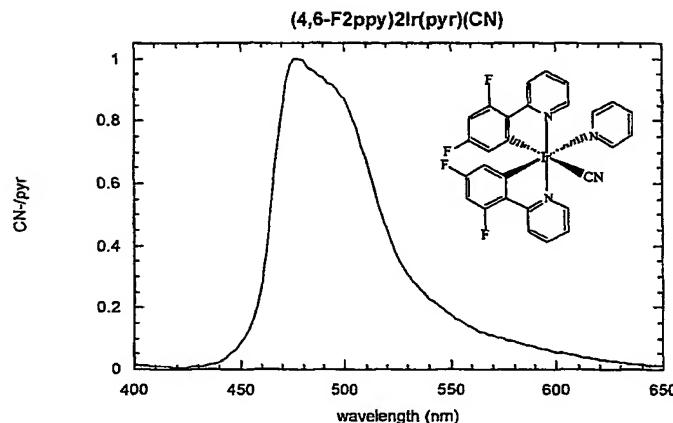
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(triphenylphosphine) cyanide.

The $[(2\text{-}(4,6\text{-difluorophenyl})\text{pyridyl})_2\text{IrCl}]_2$ complex (ca. 0.002 g) was treated with an excess of triphenylphosphine and potassium cyanide in 2 mL CH_2Cl_2 solution for 16 hours.



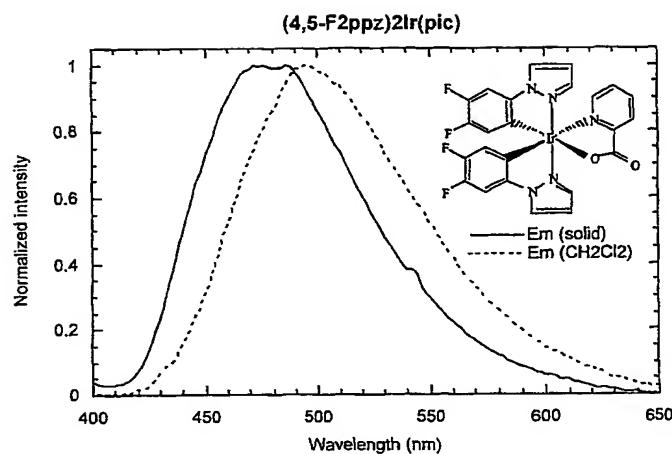
Synthesis of Iridium (III) bis(2-(4,6-difluorophenyl)pyridinato-N,C^{2'})(pyridine) cyanide.

The $[(2\text{-}(4,6\text{-difluorophenyl})\text{pyridyl})_2\text{IrCl}]_2$ complex (ca. 0.002 g) was treated with an excess of pyridine and potassium cyanide in 2 mL CH_2Cl_2 solution for 16 hours.



Synthesis of Iridium (III) bis(1-(4,5-difluorophenyl)pyrazolyl-N,C^{2'})(picolinate).

The cyclometalated Ir(III) μ -chloro bridged dimer of a general formula $C-N_2\text{Ir}(\mu-\text{Cl})_2\text{IrC}-N_2$ was synthesized by heating a mixture of $\text{IrCl}_3 \cdot n\text{H}_2\text{O}$ with 4 eq. of 1-(4,5-difluorophenyl)pyrazole) in 2-ethoxyethanol at 130°C for 16 hr. The crude product was isolated by addition of water followed by filtration and methanol wash. The crude $[(1-(4,5\text{-difluorophenyl})\text{pyrazolyl})_2\text{IrCl}]_2$ complex was treated with 2 eq of picolinic acid in refluxing 1,2-dichloroethane under inert gas atmosphere for 16 hours. After cooling down to room temperature, the solvent was removed under reduced pressure and the crude yellow product was washed with methanol to remove any unreacted picolinic acid. The crude product was flash chromatographed using a silica:dichloromethane column to yield the pure colorless $(C-N)_2\text{Ir(pic)}$ after solvent evaporation and drying.



The chemical structures of the phosphorescent organometallic compounds from Figures 7a – 7r, along with some of the ligands comprising these compounds are also shown in Figures 8a – 8d.

One example of a representative embodiment of the present invention is Iridium(III)bis(4,6-di-fluorophenyl)-pyridinato-N, C²) picolinate (FIrpic), which was used as a phosphorescent dopant in an OLED to produce efficient blue electrophosphorescence. The synthesis of FIrpic is as follows. The 2-(4,6-difluorophenyl)pyridine ligand precursor was prepared by Suzuki coupling of 4,6-difluorophenylboronic acid (Frontier Chemical) with 2-bromopyridine (Aldrich) in 1,2-dimethoxyethane using a Pd(OAc)₂/PPh₃ catalyst and K₂CO₃ base as per Synlett, 1999, 1, 45-48. Next, [(2-(4,6-difluorophenyl)pyridyl)₂IrCl]₂ was synthesized. All procedures involving IrCl₃·nH₂O or any other Ir(III) species were carried out in inert gas atmosphere in spite of the air stability of the compounds, the main concern being their oxidative stability and stability of intermediate complexes at high temperatures used in the reactions. Cyclometallated Ir(III) di-chloro bridged dimers of a general formula C-N₂Ir(mu-Cl)₂IrC-N₂ were synthesized by heating a mixture of IrCl₃·nH₂O (Next Chimica) with 4 eq. of 2-(4,6-difluorophenyl)pyridine) in 2-ethoxyethanol (Aldrich Sigma) at 130°C for 16 hr. The product was isolated by addition of water followed by filtration and methanol wash, resulting in a yield of 90%.

The general procedure for synthesizing (C-N)₂Ir(pic) complexes is as follows. The [(C-N)₂IrCl]₂ complex was treated with 2 eq. of picolinic acid (Aldrich Sigma) in refluxing 1,2-dichloroethane under inert gas atmosphere for 16 hours. After cooling down to room temperature, the solvent was removed under reduced pressure and the crude yellow product was washed with methanol to remove any unreacted picolinic acid. The crude product was flash chromatographed using a silica:dichloromethane

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column to yield ca. 75% of the pure yellow (C₆N)₂Ir(pic) after solvent evaporation and drying.

In one representative embodiment of the present invention, we demonstrate blue electrophosphorescence using energy transfer from a conductive organic host to an iridium complex with two 2-(4,6-difluoro-phenyl)pyridine as two mono-anionic, bidentate, carbon-coordination ligands (cyclometallated) and a picolinate ligand as a non-mono-anionic, bidentate, carbon-coordination ligand. See Lamansky, S., Djurovich, P., Murphy, D., Abdel-Razzaq, F., Adachi, C., Burrows, P. E., Forrest, S. R., and Thompson, M. E., J. Am. Chem. Soc., (in press). The introduction of fluorine groups, which are electron withdrawing substituents, results in an increase of the triplet exciton energy and hence a blue shift of the phosphorescence compared with that of Ir(ppy)₃. Using Iridium(III)bis(4,6-di-fluorophenyl)-pyridinato-N, C²⁻) picolinate (FIrpic), we obtained a maximum external quantum EL efficiency (η_{ext}) of (5.7±0.3)% and a luminous power efficiency (η_p) of (6.3±0.3)lm/W. To our knowledge, this is the first report of efficient blue electrophosphorescence (see Adachi, C., Baldo, M. A., Thompson, M. E., and Forrest, S. R., Material Research Society, Fall Meeting Boston, MA, 1999; Wu, Q. G., Lavigne, J. A., Tao, Y., D'Iorio, M., and Wang, S. N., Inorg. Chem., 39, 5248-5254 (2000); and Ma, Y., G., Lai, T. S., and Wu, Y., Adv. Mat., 12, 433-435 (2000)) and represents a significant improvement of the efficiencies compared with the best blue fluorescent emitters reported to date. See Grice, A. W., Bradley, D. D. C., Bernius, M. T., Inbasekaran, M., Wu, W. W., and Woo, E. P., Appl. Phys. Lett., 73, 629-931 (1998); Hosokawa, C., Higashi, H., Nakamura, H., and Kusumoto, T., Appl. Phys. Lett., 67, 3853-3855 (1995); and

Hosokawa, C., Eida, M., Matsuura, M., Fukuoka, K., Nakamura, H., and Kusumoto, T.,
Synth. Met., 91, 3-7 (1997).

Figure 1a shows photoluminescent (PL) spectra in a dilute (10^{-5} M)
5 chloroform solution of three different iridium-based phosphors, bis(2-phenylpyridinato-N,C²)iridium(acetylacetone) [ppy₂Ir(acac)] (curve c), bis (4,6-di-fluorophenyl)-pyridinato-N, C²)iridium(acetylacetone) [FIr(acac)] (curve b), and FIrpic (curve a), demonstrating a spectral shift with ligand modification. Figure 1a also shows the molecular structures of these iridium complexes: FIrpic (structure a), FIr(acac) (structure
10 b), and ppy₂Ir(acac) (structure c). The presence of the heavy metal iridium results in strong spin-orbit coupling and metal ligand charge transfer, allowing for rapid intersystem crossing of excitons into the radiative triplet manifold. See King, K. A., Spellane, P. J. and Watts, R. J., J. Am. Chem. Soc., 107, 1431-1432 (1985); and Lamansky, S.; Djurovich, P.; Murphy, D.; Abdel-Razzaq, F.; Kwong, R.; Tsyba, L; Bortz, M.; Mui, B.;
15 Bau, R.; Mark E. Thompson, M.E. *Inorganic Chemistry*, 40, 1704-1711(2001). All three of these complexes give high photoluminescent efficiencies of
 $\Phi_{pl} = 0.5\text{-}0.6$ in fluid solution. With the introduction of electron withdrawing fluorine atoms into the 4,6-positions of 2-phenylpyridine, the triplet excited state experiences a blue shift of ~40nm in the PL peak of FIr(acac), as compared with the green emitting ppy₂Ir(acac).
20 Furthermore, replacement of the acetylacetone ligand (acac) of FIr(acac) with picolinate (*i.e.*, FIrpic) resulted in an additional ~20 nm blue shift.

Organic light emitting devices (OLEDs) were grown on a glass substrate

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precoated with a ~130 nm thick indium-tin-oxide (ITO) layer with a sheet resistance of ~ $20\Omega/\square$. Prior to organic layer deposition, the substrate was degreased with solvents and cleaned for 5 min by exposure to a UV-ozone ambient, after which it was immediately loaded into the evaporation system. With a base pressure of $\sim 4 \times 10^{-8}$ Torr, the organic and metal cathode layers were grown successively without breaking vacuum using an in 5 *vacuum* mask exchange mechanism. First, a 10nm-thick copper phthalocyanine (CuPc) hole injection layer followed by a 30nm-thick 4,4'-bis[N-(1-naphthyl)-N-phenyl-amino]biphenyl (α -NPD) hole transport layer (HTL), respectively, were deposited. Next, a 10 30nm-thick light-emitting layer (EML) consisting of 6%-FIrpic doped into a 4,4'-N,N'-dicarbazole-biphenyl (CBP) host was prepared via thermal co-deposition. Finally, a 30nm-thick layer of aluminum(III)bis(2-methyl-8-quinolinato)4-phenylphenolate (BAlq) was used to transport and inject electrons into the EML. A shadow mask with rectangular 15 openings was used to define the cathode consisting of a 1 nm-thick LiF layer, followed by a 100nm-thick Al layer. After deposition, the device was encapsulated using a UV-epoxy resin under a nitrogen atmosphere with < 1ppm oxygen and water. We found 20 that carrying out the entire layer deposition process under high vacuum is crucial for obtaining high efficiencies. Given that the peak CBP triplet wavelength (see Baldo, M. A., and Forrest, S. R., Phys. Rev. B 62,10958-10966 (2000)) is $\lambda=484\text{nm}$ [(2.56 ± 0.10) eV], compared to $\lambda=475\text{nm}$ [(2.62 ± 0.10) eV] for FIrpic (see spectra in Fig. 3), the endothermic transfer process may be easily interrupted by the presence of nonradiative defect states of intermediate energy. Introduction of oxygen or water may be the source of such defects. Indeed, we have found that breaking vacuum at any point in the fabrication process and exposure to air or purified nitrogen (< 1ppm oxygen and water) results in a decrease in 25

efficiency of at least a factor of two below the values reported here. A similar ambient sensitivity is not observed for green and red electrophosphorescence OLEDs employing conventional exothermic energy transfer mechanisms.

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Figure 1b shows the electroluminescence spectra of the following OLED structure: ITO/ CuPc (10nm)/ α -NPD(30nm)/ CBP host doped with 6% FIrpic (30nm)/ BA1q (30nm)/ LiF (1nm)/ Al (100nm). The EL spectrum has a maximum at the peak wavelength of $\lambda_{\text{max}}=475\text{nm}$ and additional sub-peaks at $\lambda_{\text{sub}}=495\text{nm}$ and 540nm (arrows) which generally agrees with the PL spectral shape. The Commission Internationale de L'Eclairage (CIE) coordinates of ($x=0.16$, $y=0.29$) for a FIrpic OLED are shown in the inset of Fig. 1b along with the coordinates of green ($\text{Ir}(\text{ppy})_3$) ($x=0.28$, $y=0.62$) and red ($\text{Btp}_2\text{Ir}(\text{acac})$) ($x=0.67$, $y=0.33$) electrophosphorescence devices. Further tuning of the color purity in the blue through the molecular design of ligands still needs to occur to achieve a closer match to the National Television Standards Committee (NTSC) recommended blue for a video display (lower vertex in the inset).

10

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Figure 2 shows external electroluminescent quantum (η_{ext} : filled squares) and power (η_p : open circles) efficiencies as functions of current density for the following OLED structure: ITO/ CuPc (10nm)/ α -NPD(30nm)/ CBP host doped with 6% FIrpic (30nm)/ BA1q (30nm)/ LiF (1nm)/ Al (100nm). A maximum $\eta_{\text{ext}}=(5.7\pm0.3)\%$ (corresponding to an internal efficiency of ~30%) and a luminous power efficiency (η_p) of $(6.3\pm0.3)\text{lm/W}$ are achieved at $J=0.5\text{ mA/cm}^2$ and 0.1 mA/cm^2 , respectively. While the device shows a gradual decrease in η_{ext} with increasing current which has previously been attributed to

triplet-triplet annihilation (see Adachi, C., Baldo, M. A., and Forrest, S. R., *J. Appl. Phys.*, **87**, 8049-8055 (2000); Baldo, M. A., Adachi, C., and Forrest, S. R., *Phys. Rev. B* **62**, 10967-10977 (2000); and Adachi, C., Kwong, R. C., and Forrest, S. R., *Organic Electronics*, **2**, (2001) (in press)), a maximum luminance of $6400\text{cd}/\text{m}^2$ with $\eta_{\text{ext}}=3.0\%$ was obtained even at a high current of $J=100\text{mA}/\text{cm}^2$. These values compare favorably with $\eta_{\text{ext}}=2.4\%$ for fluorescent devices with a similar blue color emission spectrum. See Hosokawa, C., Higashi, H., Nakamura, H., and Kusumoto, T., *Appl. Phys. Lett.*, **67**, 3853-3855 (1995). The inset to Figure 2 shows an energy level diagram of triplet levels of a CBP host and a FIrpic guest. Due to the energy lineup of CBP and FIrpic triplet levels, both exothermic and endothermic transfer can be possible. Here, κ_g and κ_h are the radiative decay rates of triplets on the guest (phosphor) and host molecules, and the rates of exothermic (forward) (κ_F) and endothermic (reverse) (κ_R) energy transfers between CBP and FIrpic is also indicated. Since the triplet energy level of a CBP host (2.56 ± 0.10) eV is slightly less than that of FIrpic at (2.62 ± 0.10) eV (inset of Fig. 2), exothermic energy transfer from FIrpic to CBP is inferred. Note that the pronounced roll-off at small J is uncharacteristic of conventional electrophosphorescence. This is indicative of the sensitivity of backward energy transfer to the presence of energy dissipative pathways, reducing the efficiency via nonradiative triplet recombination when the density of triplets is too low to saturate these parasitic mechanisms. In addition, unbalanced hole and electron injection into EML can also account for the roll off.

Figure 3 shows a streak image of the transient decay of a 6%-FIrpic:CBP film (100nm thick) on a Si substrate under nitrogen pulse excitation (~500ps) at T=100K.

Two distinct decay processes, prompt and delayed phosphorescence, are demonstrated along with their photoluminescent spectra: dashed line=prompt and solid line=delayed. Also shown is the CBP phosphorescence spectrum obtained at 10K. In addition to the prompt phosphorescence of FIrpic, we observe an extremely long decay component lasting 5 for $\tau \sim 10\text{ms}$ which follows the CBP triplet lifetime. The prompt phosphorescence has a lifetime somewhat shorter than the solution phosphorescence lifetime of FIrpic ($2\ \mu\text{sec}$). Since the PL spectrum of the slow component coincides with that of FIrpic PL, this supports the conclusion that exothermic energy transfer from FIrpic to CBP occurs. The triplet state then migrates through the CBP host molecules and finally is endothermally 10 transferred back to FIrpic, resulting in the delayed phosphorescence observed. Due to the significant difference of lifetimes of the excited states, $\kappa_h \ll \kappa_g$, (κ_h and κ_g are the radiative decay rates of the triplets on the host and guest molecules, respectively), the triplet exciton decay originates from FIrpic, as desired. Here, the low intensity blue emission centered at 15 $\lambda_{\max}=400\text{nm}$ in the prompt emission spectrum is due to fluorescence of CBP, with a transient lifetime $<<100\text{ns}$ which is significantly shorter than the transient decay of FIrpic. A similar process has been observed for energy transfer from the pyrene triplet to a Ru MLCT excited state, leading to long Ru-MLCT excited state lifetimes. See Ford, W. E., Rodgers, M. A. J., *J. Phys. Chem.*, **96**, 2917-2920 (1992); Harriman, A.; Hissler, M.; Khatyr, A.; Ziessel, R. *Chem. Commun.*, 735-736 (1999).

20.

Figure 4 shows the transient photoluminescence decay characteristics of a 100nm thick 6%-FIrpic:CBP film on a Si substrate under nitrogen pulse excitation (~500ps) at T=50K, 100K, 200K and 300K. The inset to Figure 4 shows the temperature

dependence of the relative photoluminescence (PL) efficiency (η_{PL}) of FIrpic doped into CBP. After a slight enhancement of η_{PL} as temperature is increased from 50K to 200K, it once again decreases at yet higher temperatures. The transient decay characteristics are also temperature dependent. In particular, a significant decrease in the nonexponential decay time was observed at T=50K and 100K. The enhancement of η_{PL} from T=300K to 5 200K is due to the suppression of nonradiative decay of FIrpic. The decrease of η_{PL} below T~200K, however, is a signature of retardation of the endothermic process of energy transfer from CBP to FIrpic, leading to loss of the radiative triplet excitons. Since we observe no delayed component at T=300K, energy transfer from CBP to FIrpic is very 10 efficient with thermal assistance. In contrast, the PL intensity of Ir(ppy)₃:CBP shows no temperature dependence along with no evidence for such a slow component at low temperature, suggesting the absence of backward energy transfer in that low triplet energy guest system.

15 As discussed above, we have herein demonstrated efficient blue electrophosphorescence using FIrpic as the phosphor molecule. Due to the comparable energy of the phosphor triplet state relative to that of the 4,4'-N,N'-dicarbazole-biphenyl (CBP) conductive host molecule into which it is doped, the exothermic transfer of energy from phosphor to host, and subsequent endothermic transfer from host back to phosphor is clearly observed. Using this triplet energy transfer process, we force emission from the 20 higher energy, blue triplet state of the phosphor. The existence of endothermic energy transfer is confirmed by the low temperature phosphorescent spectra of CBP and FIrpic, and the appearance of a maximum in the photoluminescent intensity of CBP:FIrpic at

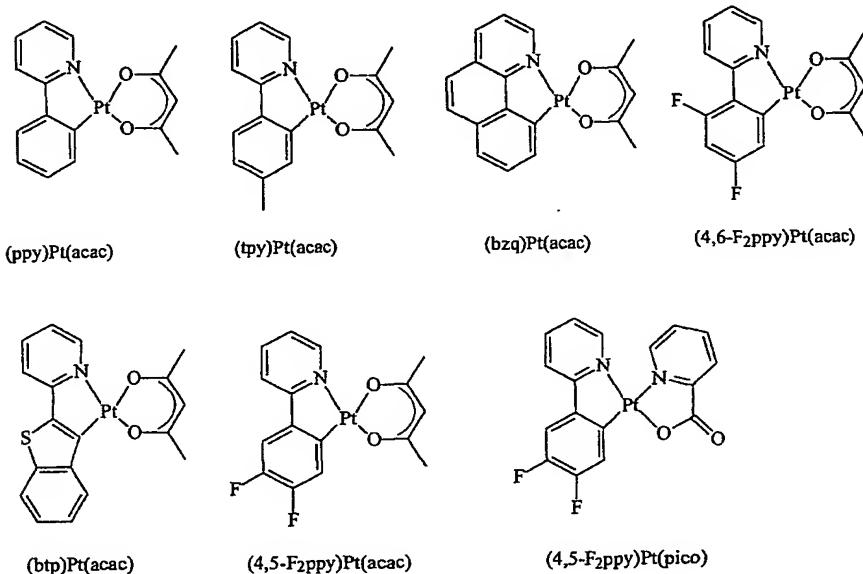
T~200K. Employing this process, a very high maximum external quantum efficiency of (5.7±0.3) % and a luminous power efficiency of (6.3±0.3)lm/W are achieved. The electroluminescent (EL) spectrum has a maximum at a wavelength of $\lambda_{\text{max}}=470\text{nm}$ with additional peaks at $\lambda_{\text{sub}}=495\text{nm}$ and 540nm, leading to the Commission Internationale de L'Eclairage (CIE) coordinates of x=0.16 and y=0.29.

Other representative embodiments of the organometallic compounds of the present invention are aimed at a new class of platinum complexes, which give efficient phosphorescence. The representative complexes have a single organometallic ligand (cyclometallated) and a bidentate coordination ligand (such as acetylacetone). Several examples of these complexes are given below along with their spectra. The emission spectra from these complexes show vibronic fine structure, consistent with strong ligand $\pi - \pi^*$ character in the phosphorescent transition. Strong ligand $\pi - \pi^*$ character is also consistent with the fact that the emission energy is strongly dependent on the identity of the ligand, as shown in the spectra shown below. Emission from these complexes results from a mixture of metal to ligand charge transfer (MLCT) and ligand based transitions.

The MLCT is critical to enhance the efficiency of intersystem crossing and phosphorescence. The emission is dominated by the cyclometallated ligand and the MLCT between the Pt ion and that ligand. The emission spectrum is only slightly affected by changing the acetylacetone ligand (acac) to a picolinic acid (pico), as shown for the (ppy)PtX complexes below. This minor shift most likely occurs due to a shift in the Pt based HOMO level by the pico ligand, leading to a red shift in the MLCT and a corresponding red shift in the emission spectrum.

One of these complexes, i.e. (2-(4,5-F₂-phenyl)pyridinato)Pt(acetylacetone), was used as a phosphorescent dopant in a polymer OLED and gave an emission spectrum identical to the photoluminescence spectrum and an external quantum efficiency of 1.3%.

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As representative electrophosphorescent compounds of the present

invention, the carbon-coordination ligand forms a cyclometallated ring that includes the organometallic carbon-metal bond and a dative bond between the metal atom and a nitrogen, sulfur or oxygen group, for example, Pt(II)-(2-phenylpyridinato-N, C²)(acetyl acetate), herein referred to as Pt(ppy)(acac) or (ppy)Pt(acac). The carbon atom that is

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bound to the metal may be present as part of a substituted or unsubstituted, saturated hydrocarbon; a substituted or unsubstituted, aromatic system, for example, phenylene or naphthalene compounds; or a substituted or unsubstituted heterocyclic system, which might include, for example, substituted or unsubstituted thiophenes, furans, pyridines and pyrroles. The group in the cyclometallated ring that forms a dative bond with the metal atom may be independently selected also to include a substituted or unsubstituted, saturated hydrocarbon; a substituted or unsubstituted, aromatic system, for example, phenylene or naphthalene compounds; or a substituted or unsubstituted heterocyclic system, which might include, for example, thiophenes, furans, pyridines and pyrroles.

10

The preparation of these representative compounds of the present invention was accomplished as follows. All procedures involving K₂PtCl₆ or any other Pt species were carried out in inert gas atmosphere in spite of the air stability of the compounds, the main concern being their oxidative stability and stability of intermediate complexes at high temperatures used in the reactions. NMR spectra were recorded on Bruker AMX 360 MHz or 500 MHz instruments unless specified otherwise. Solid probe MS spectra were taken with Hewlett Packard GC/MS instrument with electron impact ionization and model 5873 mass sensitive detector. High resolution mass spectrometry was done at Frik Chem Laboratories at Princeton University. Elemental analysis data was recorded at the Microanalysis Laboratory at the University of Illinois, Urbana-Champaine.

Pt(II) μ -chlorobridged dimers of the structure [Pt(C-N)(μ -Cl)₂Pt(C-N)] containing cyclometalated carbon, nitrogen ligands (C,N) used in the study were prepared

according to Cave G.W.V., Fanizzi F.P., Deeth R.J., Errington W., Rourke J.P.,
Organometallics 2000, 19, 1355.

Platinum(II) (2-phenylpyridinato-NC²) (acetyl acetonate) [Pt(ppy)(acac)].

5 100 mg of Pt(ppy)(μ-Cl)₂Pt(ppy) dimer, 25 mg of 2,4-pentanedione and 85 mg of
anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under
inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water
and filtration yielded crude product that after drying and flash chromatography
(silica/dichloromethane) gave bright yellow-green solid (36% yield). ¹H NMR (360 MHz,
acetone-d₆), ppm: 9.00 (d, 1H, J 5.8Hz), 8.02 (dt, 1H, J 1.6, 7.4Hz), 7.89 (d, 1H, J 7.9H),
10 7.57 (dd, 1H, J 1.6, 7.4Hz), 7.51 (dd, 1H, J 1.6, 7.9Hz), 7.32 (dt, 1H, J 1.6, 6.8Hz), 7.11
7.04 (dt, 1H, J 1.6, 7.4Hz), 5.55 (s, 1H), 1.96 (s, 3H), 1.95 (s, 3H).
See Figure 12, compound number 5. See also Figure 9(a).

15 *Platinum(II) (2-(p-tolyl)pyridinato-N,C²) (acetyl acetonate)*
[Pt(tpy)(acac)]. 100 mg of Pt(tpy)(μ-Cl)₂Pt(tpy) dimer, 25 mg of 2,4-pentanedione and 85
mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol
under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold
water and filtration yielded crude product that after drying and flash chromatography
(silica/dichloromethane) gave bright yellow-green solid (42% yield). ¹H NMR (360 MHz,
CDCl₃), ppm: 8.94 (d, 1H, J 5.9Hz), 7.74 (t, 1H, J 6.8Hz), 7.53 (d, 1H, J 7.8Hz), 7.39 (s,
1H), 7.30 (d, 1H, J 7.8Hz), 7.04 (t, 1H, J 6.8Hz), 6.88 (d, 1H, J 7.8Hz), 5.45 (s, 1H), 2.00
1.98 (s, 3H), 1.95 (s, 3H). See Figure 12, compound number 1. See also Figure

9(b).

Platinum(II) (7,8-benzoquinolinato-N,C^{3'}) (acetyl acetonate)

[Pt(bzq)(acac)]. 100 mg of Pt(bzq)(μCl)₂Pt(bzq) dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow solid (27% yield). ¹H NMR (360 MHz, acetone-d₆), ppm: 9.13 (d, 1H, J 5.4Hz), 8.25 (d, 1H, J 8.3Hz), 7.75 (m, 2H), 7.50-7.57 (m, 10 3H), 7.44 (dd, 1H, J 5.4, 5.4Hz), 5.52 (s, 1H), 2.04 (s, 6H). See Figure 12, compound number 2. See also Figure 9(c).

Platinum(II) (2-benzylpyrinato-N,C^{2'}) (acetyl acetonate) [Pt(bzpy)(acac)].

100 mg of Pt(bzpy)(μ-Cl)₂Pt(bzpy) dimer, 25 mg of 2,4-pentanedione and 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellowish green solid (20% yield). ¹H NMR (500 MHz, CDCl₃), ppm: 8.88 (d, 1H), 7.71 (t, 1H), 7.35-7.43 (m, 2H), 7.13 (t, 1H), 6.98-7.02 (m, 2H), 6.91 (t, 1H), 5.49 (s, 1H), 4.16 (s, 2H), 1.96 (s, 3H), 1.95 (s, 3H).

Platinum(II) (2-(2'-thienyl)pyridinato-N, C^{3'}) (acetyl acetonate)

[Pt(thpy)(acac)]. 100 mg of Pt(thpy)(μCl)₂Pt(thpy) dimer, 25 mg of 2,4-pentanedione and

85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright orange solid (20% yield). ^1H NMR (500 MHz, CDCl₃) ppm: 8.78 (d, 1H), 7.67 (t, 1H), 7.46 (d, 1H), 7.26 (d, 1H), 7.17 (d, 1H), 6.86 (t, 1H), 5.46 (s, 1H), 1.98 (s, 3H), 1.95 (s, 3H).

10 Platinum(II) (2-(2'-(4',5'-benzothienyl)pyridinato-*N,C*^{3'}) (acetyl acetonate)
 $[Pt(btp)(acac)]$. 100 mg of Pt(btp)(μ -Cl)₂Pt(btp) dimer, 25 mg of 2,4-pentanedione and
15 85 mg of anhydrous sodium carbonate were refluxed at 100 °C in 8 ml of 2-ethoxyethanol
under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold
water and filtration yielded crude product that after drying and flash chromatography
(silica/dichloromethane) gave orange-red solid (20% yield). ¹H NMR (360 MHz, CDCl₃),
ppm: 8.90 (d, 1H, J 5.9Hz), 8.75-8.79 (m, 1H), 7.77-7.81 (m, 1H), 7.71 (dt, 1H, J 1.5,
7.8Hz), 7.27-7.34 (m, 3H), 6.95 (dt, 1H, J 1.5, 6.8Hz), 5.54 (s, 1H), 2.08 (s, 3H), 2.01 (s,
3H). See Figure 12, compound number 3. See also Figure 9(e).

Platinum(II) (2-(4',6'-difluorophenyl)pyridinato-*N,C*²⁺) (acetyl acetonate) [Pt(4,6-F₂ppy)(acac)]. 131 mg of Pt(4,6-F₂ppy)(μ-Cl)₂Pt(4,6-F₂ppy) dimer, 43 mg of 2,4-pentanedione and 109 mg of anhydrous sodium carbonate were refluxed at 100 °C in 10 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow solid. ¹H NMR

(360 MHz, acetone-d₆, ppm: 9.06 (dt, 1H, J 1.0, 5.9Hz), 8.08-8.13 (m, 1H), 8.01 (dt, 1H, J 1.5, 8.3Hz), 7.38-7.43 (m, 1H), 7.05 (dd, 1H, J 2.4, 9.3Hz), 6.69-6.76 (m, 1H), 5.61 (s,1H), 2.01 (s, 3H), 1.99 (s, 3H). See Figure 12, compound number 4. See also Figure 9(d).

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*Platinum(II) (2-(4',5'-difluorophenyl)pyridinato-N,C^{2'}) (acetyl acetonate)
[Pt(4,5-F₂ppy)(acac)]*. 68 mg of Pt(4,5-F₂ppy)(μ-Cl)₂Pt(4,5-F₂ppy) dimer, 36 mg of 2-picolinic acid and 57 mg of anhydrous sodium carbonate were refluxed at 100 °C in 5 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature,
10 addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow solid. ¹H NMR (360 MHz, acetone-d₆, ppm: 8.99 (d, 1H, J 5.7Hz), 8.06 (dt, 1H, J 2.3, 8.0Hz), 7.90 (d, 1H, J 8.0Hz), 7.62-7.68 (m, 1H), 7.37 (tt, 1H, J 1.7, 5.7Hz), 7.20-7.25 (m, 1H), 5.58 (s,1H), 1.99 (s, 3H), 1.98 (s, 3H). See Figure 12, compound number 6. See also Figure 9(f).

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*Platinum(II) (2-(4',5'-difluorophenyl)pyridinato-N, C^{2'}) (2-picolinato)
[Pt(4,5F₂ppy)(pico)]*. 69 mg of Pt(4,5-F₂ppy)(μ-Cl)₂Pt(4,5-F₂ppy) dimer, 30 mg of 2-picolinic acid and 52 mg of anhydrous sodium carbonate were refluxed at 100 °C in 5 ml of 2-ethoxyethanol under inert gas atmosphere for 15 hours. Cooling to room temperature,
20 addition of cold water and filtration yielded crude product that after drying and flash chromatography (silica/dichloromethane) gave bright yellow solid. ¹H NMR (500 MHz, CDCl₃, ppm: 9.15 (d, 1H, J 5.6Hz), 9.05 (d, 1H, J 5.6Hz), 8.08-8.21 (m, 2H), 7.89 (td, 1H, J 1.2, 8.0Hz), 7.68-7.71 (m, 1H), 7.54 (d, 1H, J 8.0Hz), 7.32-7.36 (m,1H), 7.12-7.20

(m, 2H). See Figure 12, compound number 7. See also Figure 9(g).

*Platinum(II) (2-(4'-cyanophenyl)pyridinato-N, C^{2'}) (acetyl acetonate)
[Pt(cppy)(acac)]*

69 mg of Pt(cppy) μ -Cl)₂Pt(cfppy) dimer, 58 mg of 2-picolinoc acid and
5 52 mg of anhydrous sodium carbonate were refluxed at 100 °C in 5 ml of 2-ethoxyethanol
under inert gas atmosphere for 15 hours. Cooling to room temperature, addition of cold
water and filtration yielded crude product that after drying and flash chromatography
(silica/dichloromethane) gave bright yellow solid. ¹H NMR (360 MHz, acetone-d₆), ppm:
9.07 (dt, 1H, J 1.0, 5.9Hz), 8.14 (dt, 1H, J 1.5, 7.8Hz), 8.05 (dt, 1H, J 1.0, 8.3Hz), 7.77-7.79
10 (m, 2H), 7.46-7.50 (m, 1H), 7.43 (dd, 1H, J 1.5, 8.3Hz), 5.61 (s, 1H), 2.01 (s, 6H).

OLED preparation and testing. Polymer blend OLEDs were spun coat
from chloroform solution on patterned pre-cleaned and oxygen plasma treated indium tin
oxide (ITO) coated glass substrates and covered with vacuum-deposited aluminum(III)
15 tris(8-hydroxyquinolate) and/or Mg:Ag (10:1 weight ratio) cathode (500 Å) for the
single-layer and heterostructure architectures, respectively. Typically, 7.5 ml of a
chloroform solution contained 100 mg of PVK, 40 mg of PBD and 2.5 mg of
(45F₂ppy)Pt(acac). Chosen spincoating conditions (3000 RPM, 40 s, Specialty Coating
Systems, Inc.) led to 1300±20 Å-thick PVK:PBD:dye films as determined by ellipsometry
20 (Rudolph automatic ellipsometer equipped with a He:Ne laser). Prior to spinning, the
solutions were filtered through a 0.2 μ m filter. Tris(8-hydroxyquinoline) aluminum (III)
(Sigma-Aldrich, Inc) (Alq₃) was sublimed prior to use. All measurements on the devices
were carried out in air at -room temperature. Device current-voltage and light intensity

characteristics were measured using the LabVIEW™ program by National Instruments with a Keithley 2400 SourceMeter/2000 Multimeter coupled to a Newport 1835-C Optical Meter. Electroluminescence spectra were recorded at room temperature on a PTI QuantaMaster™ Model C-60SE spectrofluorometer.

5

Other methods known to those skilled in the art of fabricating OLEDs may be used.

Since the discovery that phosphorescent materials can be used as the emissive material in highly efficient OLEDs, there is now much interest in finding still more efficient electrophosphorescent materials and OLED structures containing such materials. High efficiency organic light emitting devices (OLEDs) using the phosphorescent dopant, *fac* tris(2-phenylpyridine)iridium ($\text{Ir}(\text{ppy})_3$), have been demonstrated using several different conducting host materials. See, for example, M. A. Baldo et al., *Nature*, vol. 395, 151 (1998); D. F. O'Brien et al., *Appl. Phys. Lett.*, vol. 74, 442 (1999); M. A. Baldo et al., *Appl. Phys. Lett.*, vol. 75, 4 (1999); T. Tsutsui et al., *Japanese J. Appl. Phys.*, Part 2, vol. 38, L1502 (1999); C. Adachi et al., *App. Phys. Lett.*, vol. 77, 904 (2000); M. J. Yang et al., *Japanese J. Appl. Phys.*, Part 2, vol. 39, L828 (2000); and C. L. Lee et al., *Appl. Phys. Lett.*, vol. 77, 2280 (2000). Since the triplet level of the metal-ligand charge transfer state of the green-emitting $\text{Ir}(\text{ppy})_3$ is between 2.5 eV and 3.0 eV, deep blue fluorophores with a peak wavelength at about 400 nm, such as 4,4'-*N,N'*-dicarbazole-biphenyl (CBP), are likely candidates as triplet energy transfer and exciton confining media. Using 6% to 10%- $\text{Ir}(\text{ppy})_3$ in CBP leads to efficient $\text{Ir}(\text{ppy})_3$

phosphorescence. In addition to the energetic resonance between the dopant and the host, the control of charge carrier injection and transport in the host layers is believed to be necessary for achieving efficient formation of radiative excitons. High electrophosphorescence efficiency has been achieved using Ir(ppy)₃ doped into CBP along with a 2,9-dimethyl-4,7-diphenyl- phenanthroline (BCP) electron transport and exciton blocking layer. M.A. Baldo et al., Appl. Phys. Lett., vol. 75, 4 (1999). In that device, the doped CBP layer was found to readily transport holes.

The compound 4,4',4"-tris(3-methylphenylphenylamino)triphenylamine ("m-MTDATA") was disclosed for use as a hole injecting material in electroluminescent OLEDs in which fluorescence was obtained from an electron transporting layer comprised of tris(8-hydroxyquinoline)aluminum ("Alq₃"). See Shirota et al., Appl. Phys. Lett., vol.65, no. 7, 807 (1994).

Four processes determine the overall efficiency of energy transfer between a host and a guest molecule: the rates of exciton relaxation on the guest and host, k_G and k_H, respectively, and the forward and reverse triplet transfer rates between guest and host, k_F and k_R, respectively. The rate equations, in the absence of exciton-formation processes, are

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$$\begin{aligned}\frac{dG}{dt} &= -k_G G - k_R G + k_p H, \\ \frac{dH}{dt} &= -k_H H - k_p H + k_R G,\end{aligned}\quad (1)$$

where G and H are the densities of guest and host triplet excitons. The solutions to equation (1) are biexponential decays of the form

$$G, H = A_1 \exp[k_1 t] + A_2 \exp[-k_2 t], \quad (2)$$

5

In various embodiments of the present invention, the materials used are (a) N,N'-diphenyl-N,N-bis(3-methylphenyl)-[1,1-biphenyl]-4,4'-diamine (TPD), (b) 2,9-dimethyl-4,7-diphenyl-1,10-phenanthroline(bathocuproine or BCP), (c) 4,4' - N, N' - dicarbazole-biphenyl (CBP), (d) Alq₃, (e) *fac* tris (2-phenylpyridine) iridium [Ir(ppy)₃]¹⁴ 10 and (f) 2,3,7,8,12,13,17,18-octaethyl-21H,23H-porphine platinum(II) (PrOEP). Of these materials, TPD and CBP are predominantly hole-transport and Alq₃; and BCP are electron-transport materials. Two phosphors are also employed as guests, Irx(ppy)₃, which emits at ~510 nm with a phosphorescent lifetime of -0.4 μs, id and PrOEP, which emits at 650 15 nm with a phosphorescent lifetime: of - 100 μs³.

20

In accordance with the present invention, faster phosphorescent rates can produce better emission results. An embodiment of the present invention concerns, among other things, the following relationship: k_phos * exp (-delta_G/kT) > k_host. The k_phos is the phosphorescence rate, that is, in the present invention greater than about 1 x 10⁵ /sec and/or 1 x 10⁶/sec. The delta_G is the difference in free energy, which in this case is substantially equal to the energy difference between the host and the guest triplets. The kT

is the thermal energy and is about 0.025 eV at typical device operating temperatures. The k_host is the host decay rate; in certain embodiments of the present invention, the k_host is equivalent to the radiative and non-radiative decay rates, equaling less than about 5×10^3 /sec. Thus, within the relationship identified, namely, $k_{\text{phos}} * \exp(\delta G/kT) > k_{\text{host}}$, if k_{phos} is about 1×10^5 /sec, kT is about 0.025 eV, and k_{host} is about 5×10^3 /sec, then δG is less than about 0.075 eV. An outer boundary for δG in this particular case can be that δG is less than about .17 eV. The outerlimit decision can be made in reference to the type of host material being used due to such factors as stability. Also, hosts with larger triplet energy levels can increase the device voltage.

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Further, in embodiments of the present invention, the overall triplet decay rate should not be too small in order to minimize other losses. In another embodiment of the present invention, k_{phos} is about 1×10^6 /sec, k_{host} is about 1×10^3 /sec and δG is less than about 0.17 eV. In another embodiment of the present invention, k_{phos} is about 1×10^5 /sec, k_{host} is about 5×10^3 /sec and δG is less than about .17 eV. In a preferred embodiment of the present invention, k_{phos} is about 1×10^6 /sec, k_{host} is about 1×10^3 /sec and δG is less than about .075 eV. Further embodiments can be based on this relationship.

20

In an embodiment of the present invention, Fig. 17 shows the molecular structures of the materials studied: (a) TPD (N,N'-diphenyl-N,N'-bis(3-merhylphenyl)-[1,1'-biphenyl]4,4'-diamine), (b) BCP (2, 9-dimethyl-4,7-diphenyl-1, 10-phenanthroline), (c) CBP (4,4'-N,N'-dicarbazole-biphenyl), (d) Alq₃ tris(8-hydroxyquinoline) aluminum,

(c) $\text{Ir}(\text{ppy})_3$, *fac* tris(2-phenylpyridine)iridium, and (f) PtOEP 2, 3, 7, 8, 12, 13, 17, 18-octaethyl-21H,23H-porphine platinum(II). Fig. 17(g) further shows the triplet dynamics in a guest host system of the present invention. Fig. 17(g) shows the rates of forward and back transfer, k_F and k_G , respectively, which are determined by the Gibb's free energy change (ΔG) and the molecular overlap. The rates of decay from the guest and host triplet states are labeled k_G and k_H , respectively.

The simplified OLED structures of the present invention are demonstrated herein for OLED structures in which the organic hole trapping material is a phosphorescent emissive guest material for which the emission is produced from a triplet excited state of an organic molecule. Thus, while the representative embodiments of the present invention are illustrated and described herein as using a phosphorescent organic material as the emissive guest material in the electron transporting host material, the full scope of the present invention can also include a fluorescent material as the emissive material in the electron transporting host material.

In Fig. 18, the structure of the electroluminescent devices is used to observe the transient response of triplet diffusion in organic host materials. Electron and hole transport layers are labeled ETL and HTL, respectively. The highest occupied molecular orbital (HOMO) obtained for each material corresponds to its ionization potential (IP). The lowest unoccupied molecular orbital (LUMO) is equal to the IP plus the optical energy gap, as determined from absorption spectra. Relative alignments of the energies in the fully assembled devices will differ from those shown. Two devices are shown: in Fig.

18(a) the host preferentially transports electrons and the exciton formation zone is at the interface between the host and α -NPD; in Fig. 18(b) the host preferentially transports holes and exciton formation is at the interface between the host and BCP. In both devices triplets are forced to diffuse through the host before reaching a phosphorescent dye.

5 Singlets formed during electrical excitation cause fluorescence within the host, thus triplet dynamics are reflected in the delay between fluorescence and phosphorescence.

For electroluminescent devices employing Alq₃ as the host, the exciton-formation zone is located at the interface of the HTL and Alq₃. In the case of HTL host materials, the exciton-formation zone is at the interface between the HTL and BCP. To study exciton diffusion, undoped layers of the host material were inserted between the exciton-formation interface and a phosphorescent layer. As shown in Fig. 18, triplets are forced to diffuse through this undoped zone before being captured by the doped luminescent, or "triplet sensing," layer. Also shown in Fig. 18 are proposed energy levels for the host and guest materials. The electron energy levels referenced to vacuum are indicated by the lowest unoccupied molecular orbital (LUMO) and the energy levels of holes are given by the highest occupied molecular orbitals (HOMO), as determined from the ionization potential for each material. It may be assumed here that the HOMO-LUMO gap is equal to the optical energy gap, as determined from absorption spectra. Under this assumption, the LUMO does not necessarily serve as the lowest conduction level for mobile electrons. Although not shown here, charge redistributions and polarization effects at the heterointerfaces are expected to alter the relative energy-level alignments when differing materials are brought into contact.

The triplet energies, measured from the highest energy peak of the PL spectra, and decay lifetimes are summarized in Table I. From the triplet energies, the free energy change ΔG on triplet transfer can be calculated for combinations of host and guest materials. Given the guest materials PtOEP and Ir(ppy)₃, it is possible to categorize several host and guest combinations based on the magnitude and sign of ΔG (see Fig. 17).
5 In one embodiment of the present invention when $\Delta G \ll 0$, the guest-host combinations where triplets on guest molecules are strongly confined include PtOEP in CBP and PtOEP in TPD. In these cases, the guest and host triplet energies are nonresonant; hence although $k_F \gg k_R$, both rates are much smaller than their resonance maxima.

10

In another embodiment of the present invention when $\Delta G < 0$, there are two examples of weak triplet confinement: PtOEP in Alq₃ and Ir(ppy)₃ in CBP. Here $k_F > k_R$, the system is close to resonance, and significant populations of both guest and host triplets exist. In yet another embodiment of the present invention when $\Delta G > 0$, in films of Ir(ppy)₃
15 in TPD the triplets are expected to reside primarily on the host with $k_R > k_F$. In another embodiment of the present invention when $\Delta G \gg 0$, Ir(ppy)₃ in Alq₃ exhibits extremely inefficient phosphorescence due to Alq₃ quenching of Ir(ppy)₃ triplets (corresponding to $k_R > k_F$), and will not be considered further.

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In another embodiment of the present invention, the efficiency of phosphorescence depends on the creation rates of triplets on the host and guest species. For example, if the bulk of excitons are formed on the guest molecular species, then efficient phosphorescence may be possible even though triplets are only weakly confined.

Therefore, to understand a particular electrophosphorescent guest-host system, we need to know the site of exciton formation. This may be determined by analysis of the phosphorescent transients. The phosphorescent material in the structures of Fig. 18 can be displaced from the exciton formation zone, forcing triplets to diffuse across several hundred angstroms of organic material prior to recombination. To measure the diffusion time, we first apply a short electrical pulse and generate singlet and triplet excitons at the ETL/HTL interface. The formation of excitons follows the current transient and is observed by measuring transient fluorescence from singlets in the host material. Then after the electrical excitation has ceased, the delay between the fluorescence and the onset of phosphorescence is measured. Either charge or triplet diffusion may be responsible for the delay, but charge diffusion can be effectively "turned off" by applying reverse bias following the excitation pulse to discharge traps and sweep out the remaining charge. Therefore, if similar delayed phosphorescence is observed in the presence and absence of reverse bias, than charge trapping on guest molecules must be significant. See Table I.

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Table I. Material triplet energies and room temperature triplet lifetimes.

Material	Triplet energy (± 0.1 eV)	Triplet lifetime
PtOEP	1.9	$1.10 \pm 10 \mu\text{s}$
Ir(ppy) ₃	2.4	$0.8 \pm 0.1 \mu\text{s}$
CBP	2.6	>1s
BCP	2.5	<10 μs
TPD	2.3	$200 \pm 50 \mu\text{s}$
Alq ₃	2.0	$25 \pm 15 \mu\text{s}$

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Since the probability of triplet transfer is proportional to the product of the electron and hole transfer probabilities, it is expected that triplet diffusion occurs at a slower rate than charge transport. However, even in cases where charge diffusion dominates the phosphorescent decay, there still may be additional triplet diffusion time.

5 For example, the delay in the triplet transport or the various species may diffuse over different distances prior to localization on a phosphorescent molecule. Hence, in those systems where delayed phosphorescence is eliminated by reverse bias, it can be concluded that charge trapping is significant, but it is possible that rapid triplet diffusion also occurs.

10 Unlike fluorescent guest-host systems, the phosphorescent systems summarized in Table II do not require energy transfer from guest to host (i.e. $\Delta G < 0$). Given minimal triplet losses in the host, the only relaxation pathway may be phosphorescence from the guest, and, as is observed in Ir(ppy)_3 in TPD, the overall electroluminescent quantum efficiency of phosphorescent OLED's with $\Delta G < 0$ can be as high as 3%. In such a system, the excitons reside primarily on the host and are eventually transferred to phosphorescent guest sites prior to emission. Although guest-host combinations with $\Delta G < 0$ generally exhibit superior performance by minimizing losses at the host, systems with $\Delta G < 0$ may be useful for high energy triplet emitters such as blue phosphors.

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Similar to Ir(ppy)_3 in TPD, triplet diffusion from host to guest is observed for PtOEP in Alq_3 . From transient analyses of exciton transport in Alq_3 , From transient analyses of exciton transport in Alq_3 , it is likely that the transport is dispersive with

behavior similar to charge transport. Nevertheless, in approximating it as a nondispersive system, we obtain a diffusion coefficient of $D = (8 \pm 5) \times 10^{-8} \text{ cm}^2/\text{s}$ and triplet lifetime of $\tau = (25 \pm 15) \mu\text{s}$.

5 TABLE II. The electrophosphorescent quantum efficiencies and several properties of a range of material combinations.

Guest (lifetime)	Host	ΔG ($\pm 0.1 \text{ eV}$)	Host lifetime	Emission lifetime (μs)	Trapping on guest	El quantum efficiency
PtOEP ($110 \pm 10 \mu\text{s}$)	CBP	-0.7	>1 s	80 ± 5	Yes	6%
	Ir(ppy) ₃	-0.5	<0.1 μs	80 ± 5	?	3%
	TPD	-0.4	$200 \pm 50 \mu\text{s}$	80 ± 5	Yes	3%
	Alq ₃	-0.1	$25 \pm 15 \mu\text{s}$	40 ± 5	No	3%
Ir(ppy) ₃ ($0.8 \pm 0.1 \mu\text{s}$)	CBP	-0.2	>1 s	0.4 ± 0.50	Yes	8%
	TPD	+0.1	$200 \pm 50 \mu\text{s}$	15 ± 2	No	3%
	Alq ₃	+0.4	$25 \pm 15 \mu\text{s}$	<0.1	?	<0.1%

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All the guest-host systems employing the green phosphor Ir(ppy)₃ exhibit weak triplet confinement on the phosphorescent guest, i.e., $\Delta G \sim 0$. Indeed, reverse transfer from Ir(ppy)₃ to CBP is undoubtedly responsible for some losses in luminescence efficiency and the decrease in phosphorescent lifetime from ~ 0.8 to $\sim 0.4 \mu\text{s}$. In spite of this, external quantum efficiencies as high as 8% have been obtained from Ir(ppy)₃ doped in CBP.⁴ As confirmed by the transient studies here, these efficiencies are possible

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because a majority of excitons are formed directly on Ir(ppy)₃ following charge trapping. The deep HOMO level of CBP, in particular, appears to encourage hole trapping on phosphorescent guest. But there remains significant room for improvement, and at least a further doubling of phosphorescent efficiency should be possible given the right host material.

The deposition techniques for any of the above-listed layers and materials are well-known in the art. For example, a representative method of depositing the OLED layers is by thermal evaporation or spin coating if a polymer LED is used; a representative method of depositing metal layers is by thermal or electron-beam evaporation; and a representative method of depositing indium tin oxide is by electron-beam evaporation or sputtering.

The present invention may also be used to provide stable, efficient, high brightness, monochromatic, multicolor, or full-color, flat panel displays of any size. The images created on such displays could be text or illustrations in full-color, in any resolution depending on the size of the individual OLEDs. Display devices of the present invention are therefore appropriate for an extremely wide variety of applications including billboards and signs, computer monitors, and telecommunications devices such as telephones, televisions, large area wall screens, theater screens and stadium screens. The structures described herein are included, for example, in a plurality of pixels in a light emitting device or as part of a single-pixel, flat panel backlight device. Moreover, the structures described herein may be used as part of a laser device. Because of the

exceptionally high luminous efficiencies that are possible for phosphorescent-based OLEDs, as compared with OLEDs generally, and especially with respect to conventional inorganic LEDs, the phosphorescent-based OLEDs of the present invention may also be used as a light source for illumination applications. Such light sources could replace conventional incandescent or fluorescent lamps for certain types of illumination applications. Such phosphorescence-based OLEDs could be used, for example, in large planar light sources that produce illumination of a desired color.

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If the host material is an electron transporter, then it is used as an electron transport layer (ETL) and employed in the structure of Fig. 18(a). Of the host materials used, Alq₃ and BCP films may serve as an ETL; the remaining materials are predominantly hole conductors and can be used in hole-transport layers (HTL's). For these materials, a wide-energy-gap hole and exciton-blocking material are required to contain the excitations within the HTL. For this purpose, we used BCP in structures shown in Fig. 18(b). BCP can conduct electrons but blocks holes from entering the ETL.

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The devices were fabricated by thermal evaporation of the source materials under high vacuum ($\approx 10^{-6}$ Torr). In successive evaporations, a hole-transport material was deposited on a precleaned glass substrate coated with ITO. This was followed by deposition of the host material. Since the host was also an HTL, a 120 Å thick BCP blocking layer was employed. All devices employed an Alq₃ ETL to separate the emissive region from the 1000-Å- thick 20:1 Mg:Ag cathode, thereby positioning the luminescent region more favorably within the microcavity created by the metal cathode. The devices

were completed by depositing a 500-Å-thick layer of Ag to protect the Mg-Ag cathode from oxidation. Metal depositions were defined by a shadow mask with an array of 1-mm-diameter openings.

5 Transient measurements were obtained by applying a narrow (200-ns) voltage pulse to the device under test and coupling the emission into a streak camera. This pulse width is chosen to be less than the radiative rate of the phosphors and larger than the charging time of the OLED, which for a 50- Ω load and a typical capacitance of 1 nF is about 50 ns. In some cases, the sample was placed under -10 V reverse bias following the
10 electrical pulse. External EL quantum efficiency measurements were made by placing the completed OLED directly onto the surface of a calibrated silicon photodetector and capturing every photon emitted in the forward (viewing) direction. All measurements were performed in air, except for those at low-temperature which were performed in an evacuated closed-cycle refrigerator.

15 The host materials, TPD, CBP, and Alq₃ are fluorescent and possess small or negligible phosphorescence at room temperature due to competing thermally activated nonradiative decay processes. In addition to intramolecular pathways, these nonradiative processes include triplet diffusion to defect sites followed by dissipative transitions.
20 Reducing the temperature slows the rate of phonon-assisted decay and triplet diffusion, and the phosphorescent PL spectra for TPD, CBP, and BCP at T= 10 K are shown in Fig.
19 together with the room-temperature spectra of PtOEP and Ir(ppy)₃. After extended sampling, it was possible to obtain the room-temperature phosphorescent spectra and

lifetimes for TPD and CBP. These measurements were possible because the triplet lifetimes of these materials are relatively long at room temperature: $200 \pm 50 \mu\text{s}$ and $>1 \text{ s}$, respectively. In fact, under ultraviolet excitation, weak orange CBP phosphorescence is visible to the naked eye at room temperature. In contrast, the triplet lifetime of BCP decreases rapidly as temperature increases from $\sim 1 \text{ s}$ at 10 K to $<10 \mu\text{s}$ at room temperature, although we note that short triplet lifetimes may be dominated by energy transfer to physical or chemical defects.

No phosphorescence was observed from Alq_3 even at temperatures as low as $\sim 10 \text{ K}$. In their study of hydroxyquinoline complexes, Ballardini *et al.*, were similarly unsuccessful in observing phosphorescent emission from Alq_3 , although they could observe the phosphorescent spectra of hydroxyquinoline complexes as Pb, Bi, Rh, Ir, and Pt. These latter materials all show triplet emission at 590–650 nm and while it is not certain that the triplet energy of Alq_3 also lies within this range, it seems likely that it is significantly red-shifted from the spectra of the other host materials in Fig. 19.

In Fig. 20, the transient responses of four electrophosphorescent material systems are shown. The device in Fig. 20(a) consists of a 600-A-thick Alq_3 diffusion layer and a phosphorescent sensing layer of 8% PtOEP doped in Alq_3 (8% PtOEP: Alq_3). The PtOEP: Alq_3 transients clearly exhibit delayed phosphorescence due to triplet diffusion in Alq_3 and also show minimal change when reverse bias is applied to the empty traps. However, in Fig. 20(b) we observe that a similar structure using a 8% PtLOEP:CEP emission layer fails to show delayed phosphorescence when reverse bias is applied. When

reverse bias is absent, the decay lifetime is increased due to the transport of electrons across the 400-Å-thick diffusion layer, see Fig. 20(d). Here, the observed lifetime of Ir(ppy)₃ is ~15 μs, significantly longer than its natural radiative decay of 1 μs. Taken together with the apparent absence of charge trapping on Ir(ppy)₃ in TPD to Ir(ppy)₃ is the rate limiting step in Ir(ppy)₃ phosphorescence.

All systems exhibit delayed phosphorescence in the absence of reverse bias; however, only (a) PtOEP: Alq₃ and (d) Ir(ppy)₃ =TPD delayed phosphorescence in the presence of a strong negative bias. Thus, we conclude that triplet energy transfer is present in these systems but that the other systems. (b) PtOEP:CBP and (c) Ir(ppy)₃: CBP, are dominated by charge trapping and exciton formation directly upon the phosphorescent molecular indeed, the deep HOMO level of CBP makes hole trapping on the guest likely when it is used as a host. In Sec. VII of the article entitled *Transient analysis of organic electrophosphorescence: I. Transient analysis of triplet energy transfer*, M. A. Baldo and S.R. Forrest, Physical Review B, vol. 62, no. 16, (2000), which is hereby incorporated by reference in its entirety, we discuss the relative merits of trapping and energy transfer as mechanisms for generating very high efficiency phosphorescent emission in OLEDs. But in the remainder of this and the following section we concentrate on those systems exhibiting energy transfer: notably, PtOEP: Alq₃ and Ir(ppy)₃: TPD.

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In Fig. 20(d), the peak in the delayed phosphorescence of Ir(ppy)₃: TPD occurs over 100 μs after excitation. If we examine the transient of an Ir(ppy)₃: TPD device where there is no layer separating the exciton-formation interface from the luminescent

zones, we find that delayed phosphorescence is absent and that the observed lifetime after electrical excitation is $15 \pm 2 \mu\text{s}$ (see Fig. 21(a)). Except for an initial peak containing some TPD fluorescence the decay is monoexponential and completely comprised of Ir(ppy)₃ emission. The PL decay of 10% Ir(ppy)₃:TPD also exhibits long-lived differs in that the initial peak is larger than that found in the EL decay, and no emission is observed from TPD.

The data of Figs. 20(d), 21(a), and 21(b) are to be compared with the natural phosphorescent lifetime of Ir(ppy)₃, which is only $1 \mu\text{s}$. As expected from the relative phosphorescent spectra and lifetimes of TPD and Ir(ppy)₃, these data can be explained by triplets residing for extended periods on TPD molecules. The rate of forward transfer (k_F) is slow ($\sim 15 \mu\text{s}$) and dominates the phosphorescent lifetime of Ir(ppy)₃ in TPD. We note that because the observed $15 \mu\text{s}$ Ir(ppy)₃ EL decay is also observed in the PL response, there must be significant populations of triplets in TPD after photoexcitation, i.e., $k_R \gg k_F$. The EL quantum efficiency of Ir(ppy)₃ in TPD is $\eta \sim 3\%$, providing evidence that efficient electrophosphorescence is possible even if it is energetically unfavorable for triplets to reside for an extended duration on the phosphor.

In Fig. 20, the transient response of four archetypal phosphorescent guest-host systems is shown. The PtOEP transients were recorded at $\lambda = 650 \pm 10 \text{ nm}$ and the Ir(ppy)₃ transients at $\lambda = 530 \pm 30 \text{ nm}$. The initial peaks in the transient decays are host fluorescence at the wave lengths of interest they mark the formation of singlet excitons. Triplet energy transfer is demonstrated in (a) by PtOEP: Alq₃. These transients exhibit

strong delayed phosphorescence due to triplet diffusion in Alq₃ and also show minimal change when reverse bias is applied to empty traps. This device had a 600-A-tick Alq₃ diffusion layer and a phosphorescent sensing layer of 8% PtOEP in Alq₃. However, in (b)

5 we observe that PtOEP: CBP fails to show delayed phosphorescence when reverse bias is applied, indicating that charge trapping on PtOEP is significant. This device had a 400-A-thick CBP diffusion layer and a phosphorescent sensing layer of 8% PtOEP in CBP.

Similarly, in (c) the transient response of Ir(ppy)₃:CBP also exhibits charge trapping on Ir(ppy)₃. This device had a 500-A-thick CBP diffusion layer and a phosphorescent sensing layer of 6% Ir(ppy)₃ in CBP. Energy transfer to Ir(ppy)₃ is observed in Ir(ppy)₃: tpd (d).

10 Here, the observed lifetime of Ir(ppy)₃ is 15 μ s significantly longer than its natural radiative lifetime of 1 μ s. Taken together with the apparent absence of charge trapping on Ir(ppy)₃ in TPD, this long lifetime indicates that energy transfer from TPD to Ir(ppy)₃ might be the rate limiting step in Ir(ppy)₃ phosphorescence. This device had a 200-A-thick TPD diffusion layer and a phosphorescent sensing layer of 6% Ir(ppy)₃ in TPD. Note that 15 the intensity of each transient measurement is arbitrary.

Fig. 21 shows (a) the electroluminescent response of 8% Ir(ppy)₃ in TPD.

The device contains no diffusion layer yet the lifetime of Ir(ppy)₃ TPD is significantly longer (15 μ s) than the natural radiative lifetime of Ir(ppy)₃ (\sim 1 μ s). The initial peaks in

20 the response is principally due to fluorescence from TPD at T = 292 K and T = 200 K.

The lifetime increases at low temperatures, consistent with a thermally activated process.

However, unlike the EL response, the initial transient in the photoluminescent response is comprised entirely of emission from photoexcited Ir(ppy)₃.

Fig. 22 shows the normalized, phosphorescent transients for PtOEP in Alq₃ recorded at 650 nm for diffusion distances of (a) 200 Å, (b) 400 Å, (c) 600 Å and (d) 300 Å. Also shown are the calculated transients (smooth curves) based on nondispersing diffusion of triplets given diffusion coefficient of D= (8 ± 5) x 10 μs cm²/s, and a triplet exciton lifetime in Alq₃ of r = 25 ± 15 μs.

Fig. 23 shows the exciton current incident in the phosphorescent zone for diffusion distances (a) 200 Å, (b) 400 Å, (c) 600 Å and (d) 800 Å (points). The current is calculated (points) by deconvolving the phosphorescent decay of PtOEP from the traces in Fig. 22. Also shown are the best fits assuming nondispersing behavior, assuming that the concentration of exciton formation decreases exponentially with distance from the HTL/ETL interface with a characteristic length of L=120 Å (Ref. 21). The smooth curves in Fig. 22 are calculated from these fits by convolving them with the PtOEP phosphorescent decay.

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Previous work has demonstrated the existence of triplet diffusion in Alq₃, and in Figs. 22 and 23 we study the behavior of diffusing triplets as a function of time and distance. Layers consisting of 8% PtOEP: Alq₃ are used to detect the triplets via electrophorescence; but in contrast to the other devices grown in this work, here we vary the thickness of the Alq₃ spacer layer and observe the changes in the phosphorescent decay transient. Figure 22 shows the normalized transient responses of PtOEP phosphorescence at λ_c= 650 nm for OLEDs with spacer layers of thickness (a) 200 Å, (b) 400 Å, (c) 600 Å,

and (d) 800 Å. All traces exhibit delayed phosphorescence under reverse bias, demonstrating the presence of triplet diffusion.

These delayed responses are understood as convolutions of the rate of 5 triplet arrival at the phosphorescent sensing layer with the phosphorescent decay of PtOEP. By deconvolving the phosphorescent lifetime of PtOEP from the observed decay, we can therefore extract the triplet exciton current entering the phosphorescent sensing layer of each device. The exciton current is calculated from the data of Fig. 22 by subtracting the initial fluorescent spikes, smoothing and then deconvolving the 10 phosphorescent decay of PtOEP.

The triplet-exciton current, shown by the data points in Fig. 23, can be fitted to the diffusion equation:

$$\frac{d\phi}{dt} = -\frac{\phi}{\tau} D \frac{d^2\phi}{dx^2} \quad (8)$$

As shown by the solid lines in Fig. 23, this fit is used to obtain values for 15 the lifetime (τ) of Alq₃ triplets and also their diffusion constant ($D\tau$). Finally, as a check, the predicted exciton currents are reconvolved with the PtOEP phosphorescent decay and compared to the measured transients of Fig. 22 (solid lines). For these fits, we assume that the exciton formation zone is the same in each device and that the exciton concentration

decreases exponentially with distance from the HTL interface with a characteristic length of $L \sim 120 \text{ \AA}$. The spikes at $r \sim 0$ observed in the deconvolved exciton current are due to excess triplets formed within the phosphorescent zone and may indicate the presence of residual charge trapping.

From both Figs. 22 and 23, we find that the simple theory provides a reasonable approximation to the observed transient decays of the PtOEP: Alq₃ system. Nevertheless, given a single value of $D r$, it is impossible to reproduce both the sharp initial increases in the phosphorescent transients and also their long tails. Thus, similar to charge transport, the data provide evidence for dispersive exciton transport, either due to the presence of exciton traps or a distribution in the diffusion coefficient arising from variation in molecular conformations within the amorphous Alq₃ film.

The data fall into two regimes: for short diffusion distances, $D r$ determines the observed exciton currents, and at longer distances the currents are limited by the exciton lifetime r . From fits to devices with a spacer layer thickness of 200 \AA or 400 \AA we obtain a diffusion coefficient of $D r = (8 \text{ 200 or 400 \AA} \pm 5) \times 10^{-8} \text{ cm}^{2/\nu}$, and from fits to the 600- and 800 - \AA devices, we obtain an exciton lifetime of $r = 25 \pm 15 \mu\text{s}$ taken together to yield a diffusion length of $L_d = 140 \pm 90 \text{ \AA}$. This is less than the length calculated previously² at $J = 6.5 \text{ mA/cm}^2$, however, the current densities applied during the 200m excitation pulses are significantly higher ($J \sim 2500 \text{ mA/cm}^2$). It is expected that the increased triplet-triplet annihilation and triplet-charge carrier quenching at high injection levels causes the observed reduction in diffusion length.

Previously, the single exciton diffusion coefficient in Alq₃ was measured to be D_s = (1.2 ± 0.8) X 10⁻⁵ cm²/s (Ref. 22) and D_s = 2.6 X 10⁻⁴ cm²/s (Ref. 23). The diffusion coefficient of triplets is typically lower than that of singlets since both the donor and acceptor transitions are disallowed.

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Another embodiment of the present invention includes an organic light emitting layer wherein the energy difference between the lowest triplet excited state of the guest material and a corresponding relaxed state of the guest material has a corresponding wavelength of about 420 nm to 480 nm for blue light emission.

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Another embodiment of the present invention includes an organic light emitting layer wherein the energy difference between the lowest triplet excited state of the guest material and a corresponding relaxed state of the guest material has a corresponding wavelength of about 480 nm - 510 nm for aqua-blue light emission.

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Another embodiment of the present invention includes an organic light emitting layer wherein the host material has a bandgap with an energy difference corresponding to about 470 nm and the guest material has a lowest triplet excited state at an energy level at about 450 nm.

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Another embodiment of the present invention includes an organic light emitting layer wherein a plurality of guest materials are dispersed in the host material.

According to the present invention, the peak in the phosphorescent blue emission is preferably produced at a wavelength of ≤ 470 nm, more preferably at a wavelength of ~ 450 nm.

Those with skill in the art may recognize various modifications to the embodiments of the invention described and illustrated herein. Such modifications are intended to be covered by the spirit and scope of the present invention. That is, while the invention has been described in detail with reference to certain embodiments, it will be recognized by those skilled in the art that there are other embodiments of the invention within the spirit and scope of the claims.

WHAT IS CLAIMED IS:

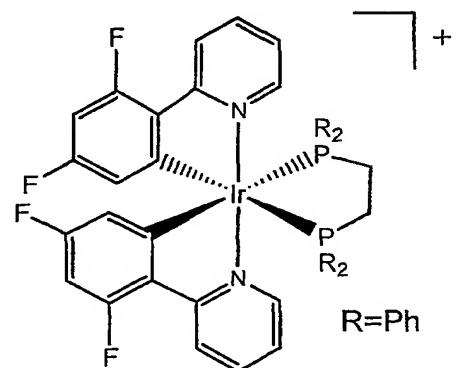
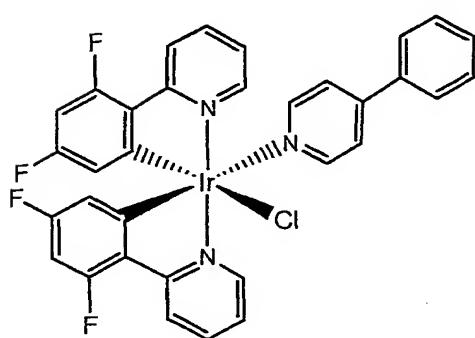
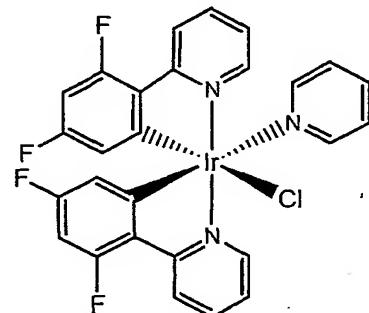
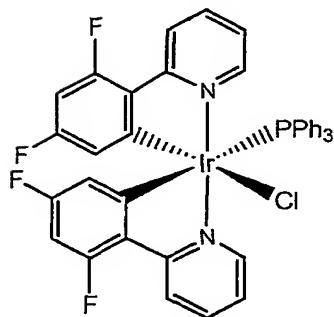
1. An organic light emitting device including an emissive layer comprising an organometallic compound comprised of:
a heavy transition metal that produces an efficient phosphorescent emission at room temperature from a mixture of metal-to-ligand charge transfer and $\pi - \pi^*$ ligand states;
at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the at least one mono-anionic, bidentate, carbon-coordination ligand is substituted with at least one of an electron donating substituent and an electron withdrawing substituent, wherein the at least one of an electron donating substituent and an electron withdrawing substituent shifts the emission, relative to an un-substituted mono-anionic, bidentate, carbon-coordination ligand, to either the blue, green or red region of the visible spectrum;
and
at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the at least one non-mono-anionic, bidentate, carbon-coordination ligand causes the emission to have a well defined vibronic structure.
2. The organic light emitting device of claim 1, wherein the heavy transition metal is selected from the group consisting of Os, Ir, Pt and Au.

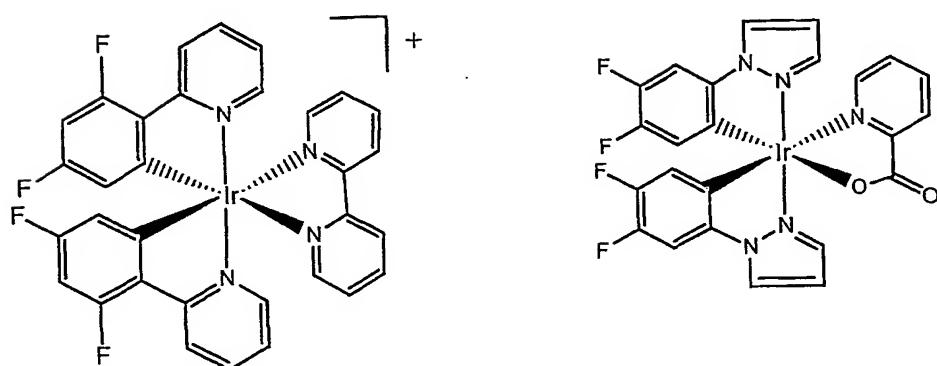
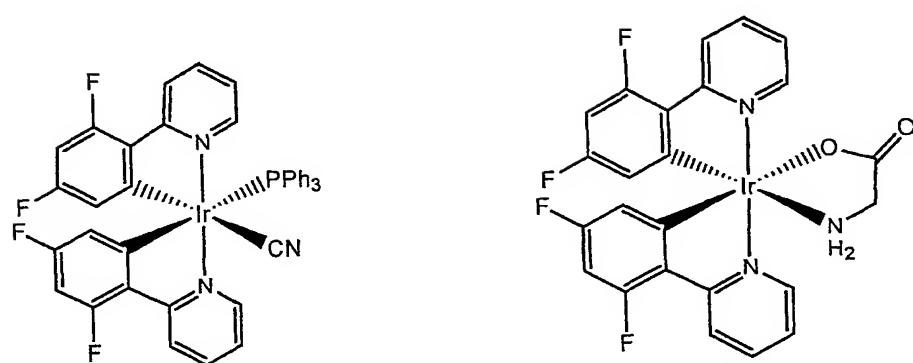
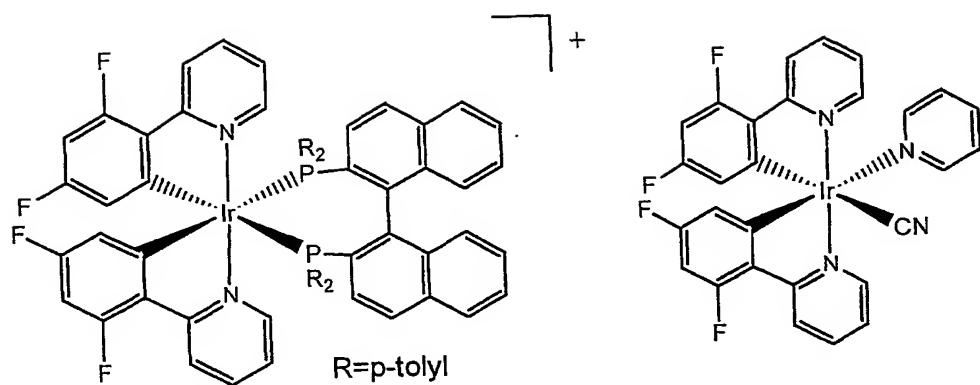
3. An organic light emitting device including an emissive layer comprising an organometallic compound comprised of:

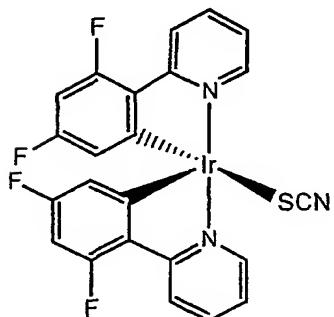
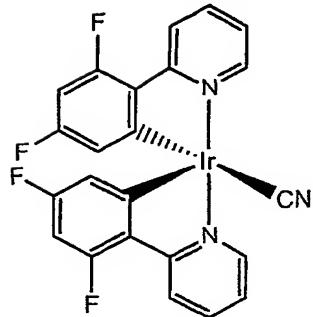
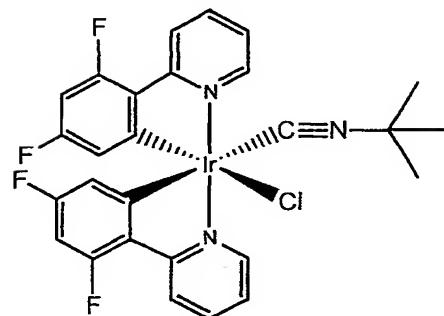
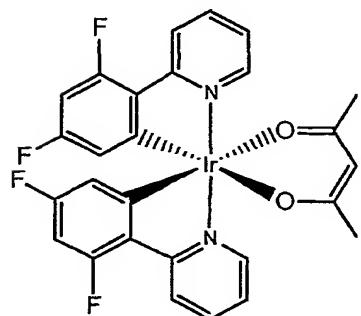
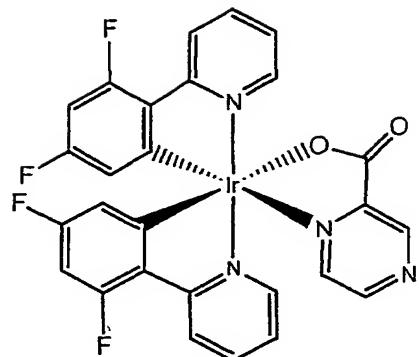
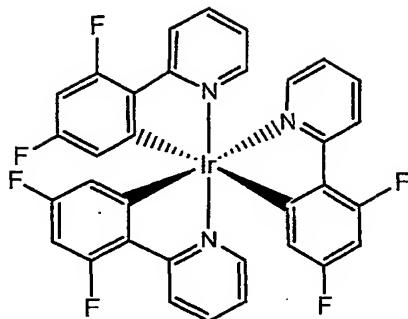
a heavy transition metal;

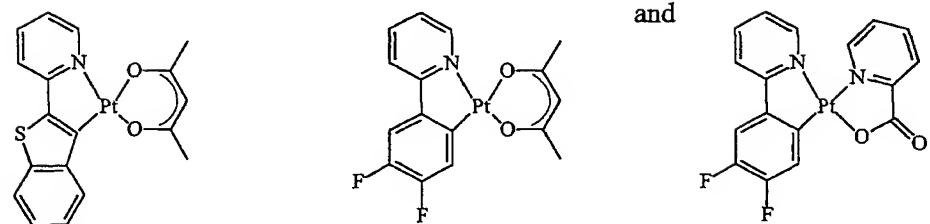
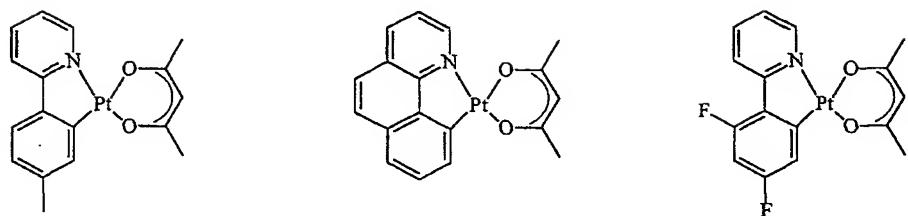
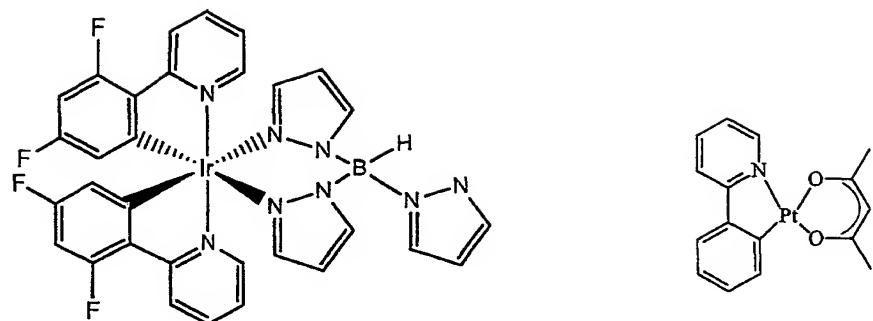
at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal; and

at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the organometallic compound has a chemical structure represented by a formula selected from the group consisting of:

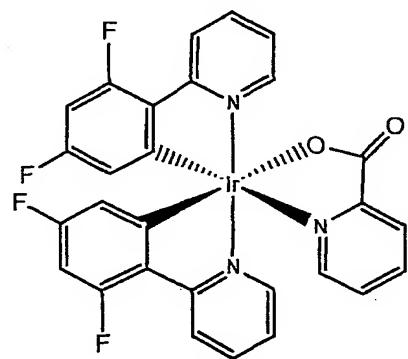




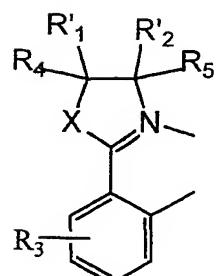
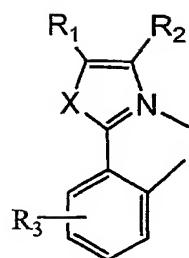
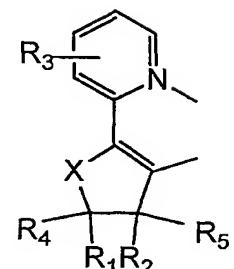
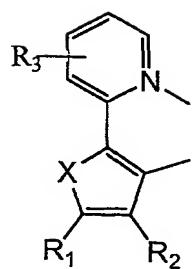
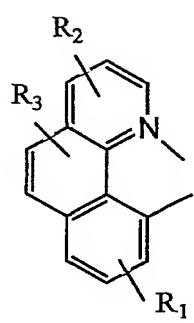
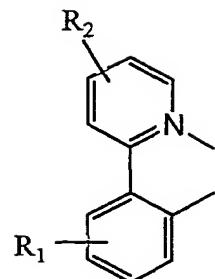
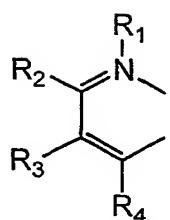


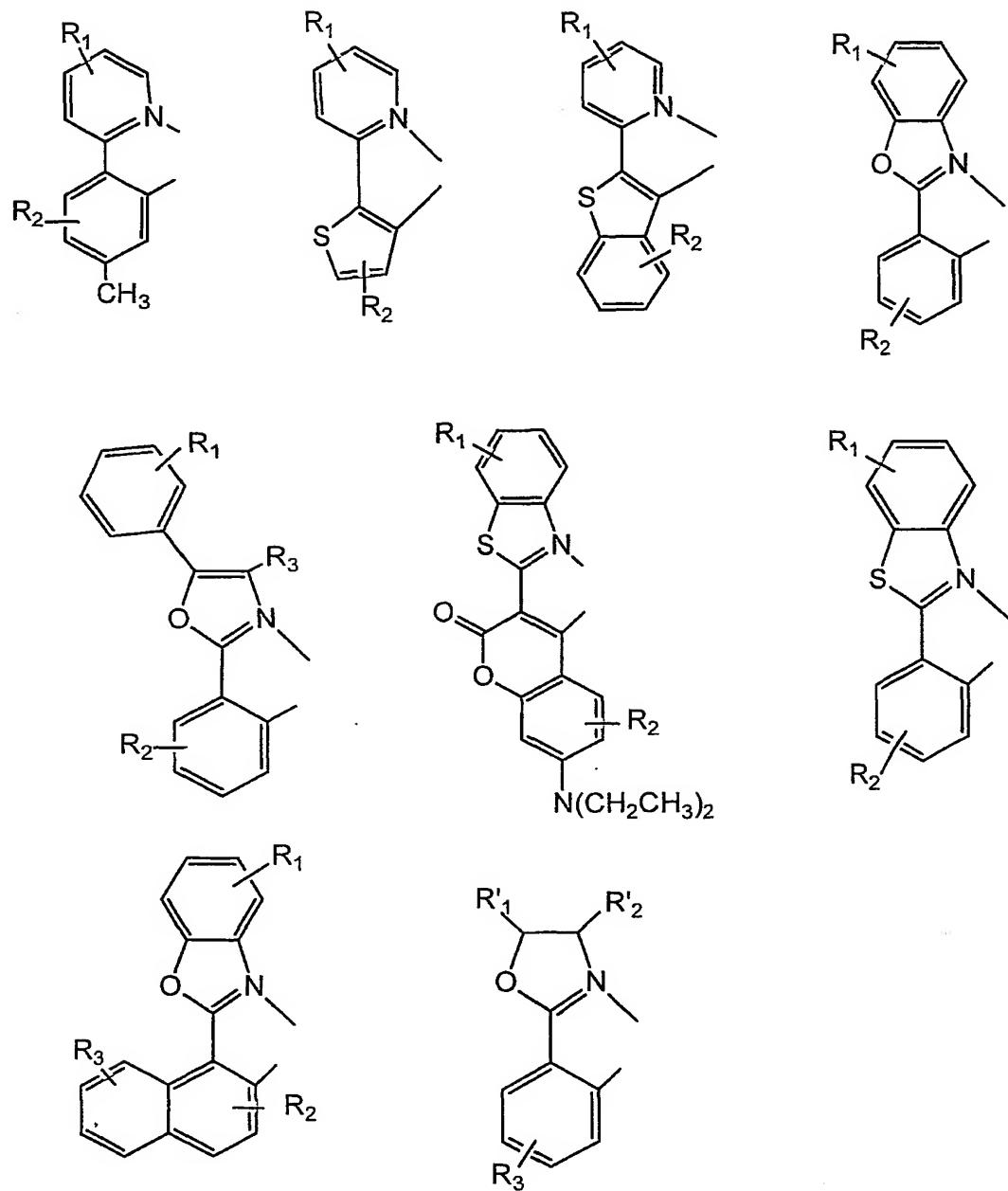


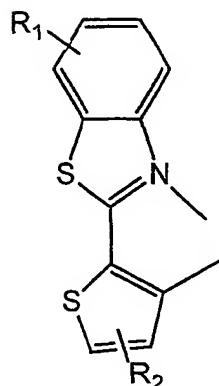
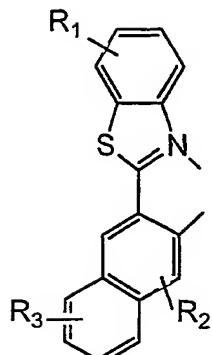
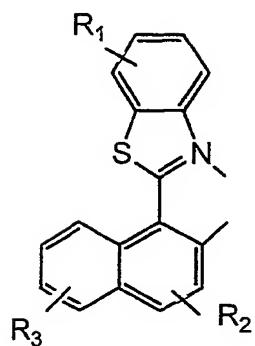
4. An organic light emitting device including an emissive layer comprising an organometallic compound comprised of:
 - a heavy transition metal;
 - at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal; and
 - at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the organometallic compound has a chemical structure represented by the following formula:



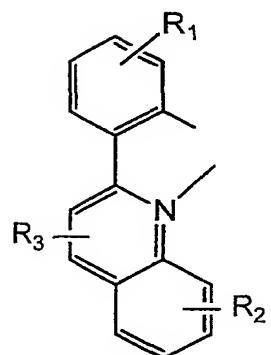
5. The organic light emitting device of claim 1, wherein the at least one mono-anionic, bidentate, carbon-coordination ligand is selected from the group consisting of :





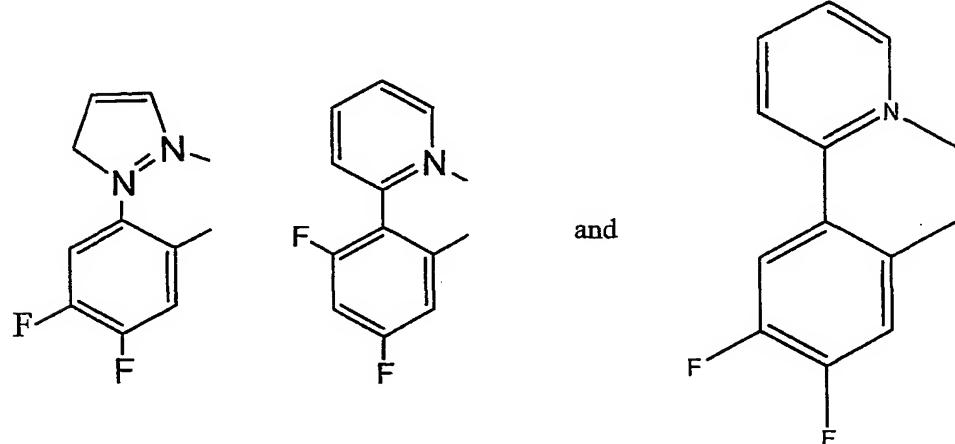


and

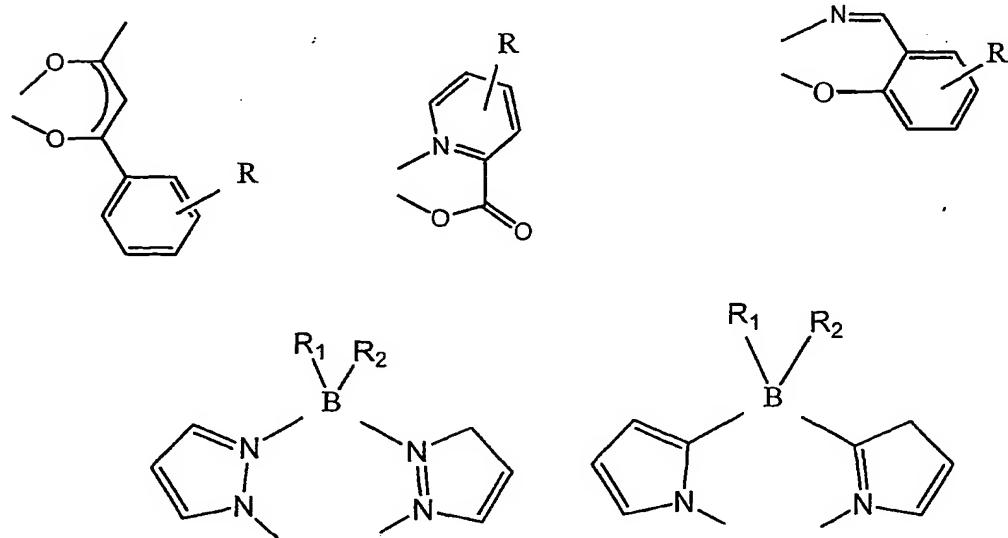


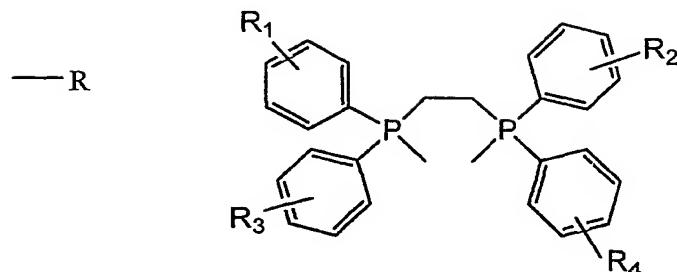
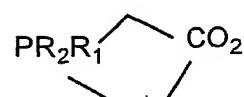
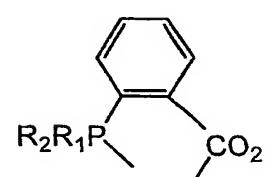
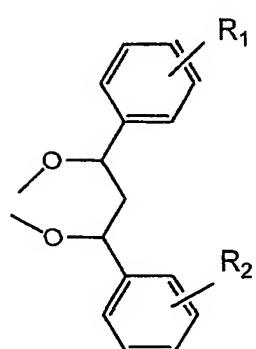
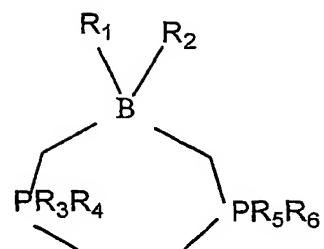
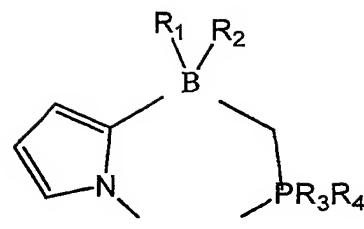
wherein X = S, O, NR; and R₁, R₂, R₃, R₄ and R₅ are, independently, hydrogen, halogen, alkyl, aryl or arylene; and R'₁ and R'₂ may, in combination, be aryl.

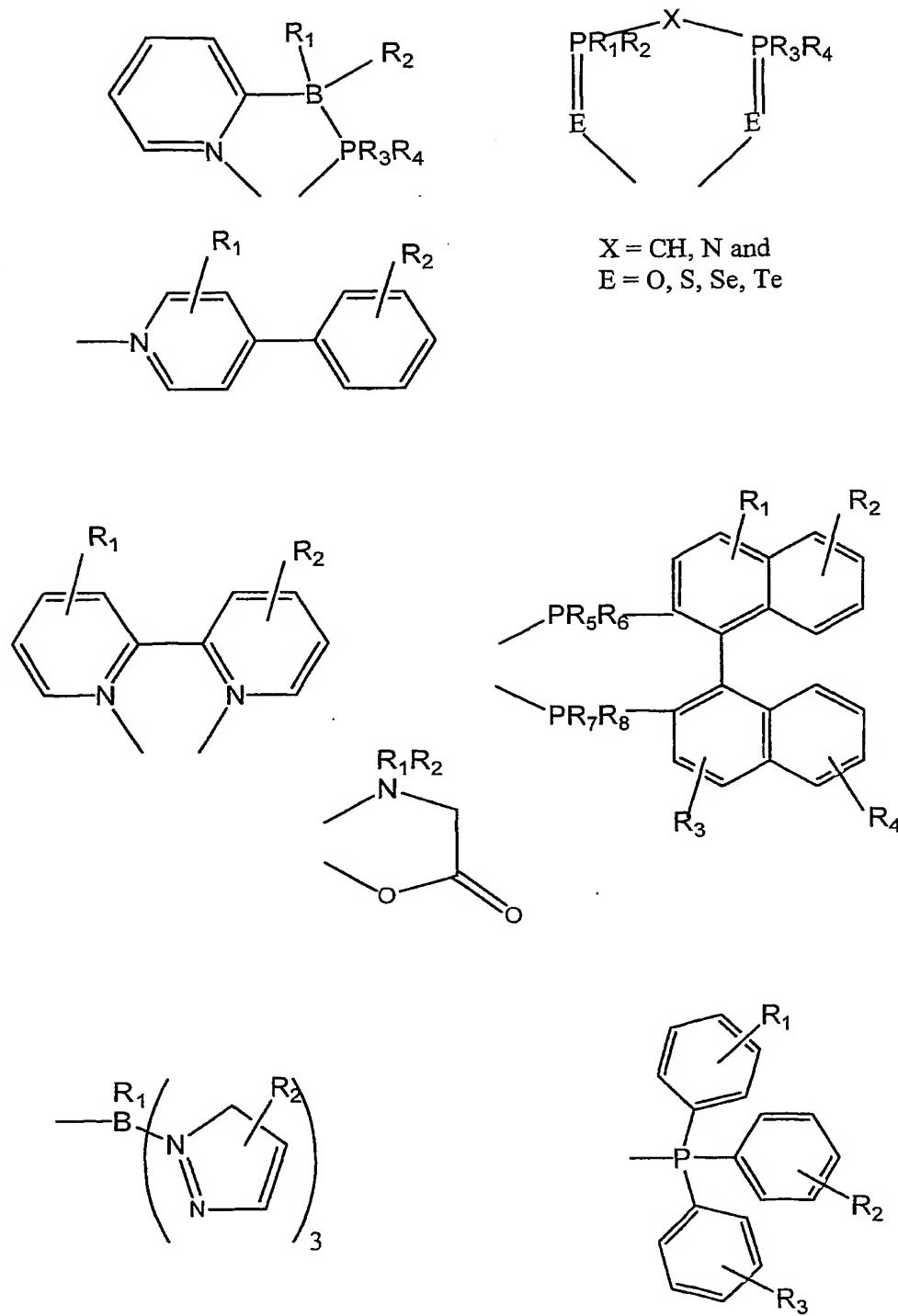
6. The organic light emitting device of claim 1, wherein the at least one mono-anionic, bidentate, carbon-coordination ligand is selected from the group consisting of:



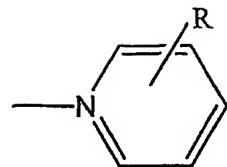
7. The organic light emitting device of claim 1, wherein the at least one non-mono-anionic, bidentate, carbon-coordination ligand is selected from the group consisting of:





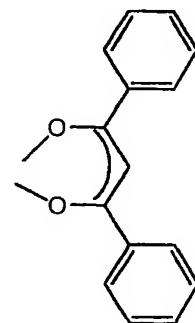
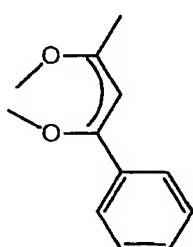
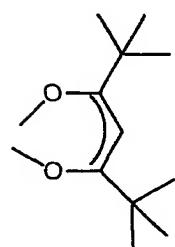
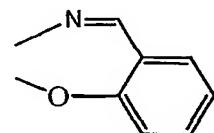
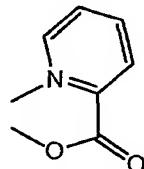
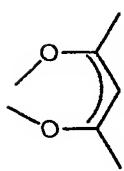


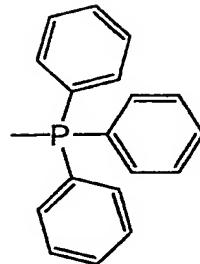
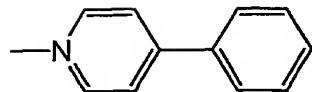
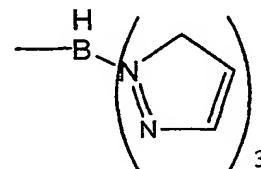
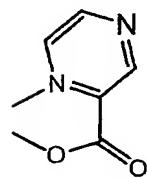
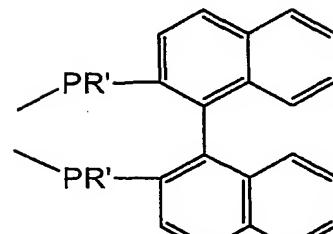
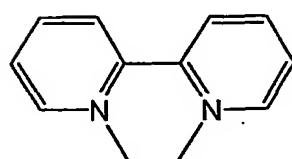
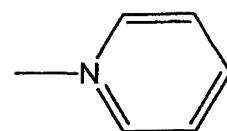
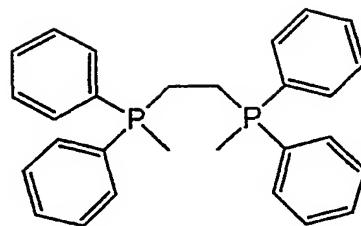
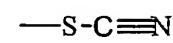
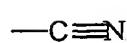
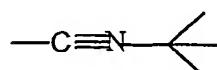
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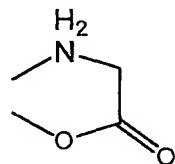
wherein R, R₁, R₂, R₃, R₄, R₅, R₆, R₇ and R₈ are, independently, hydrogen, halogen, alkyl or aryl.

8. The organic light emitting device of claim 1, wherein the at least one non-mono-anionic, bidentate, carbon-coordination ligand is selected from the group consisting of :

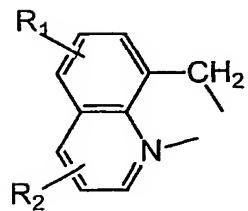
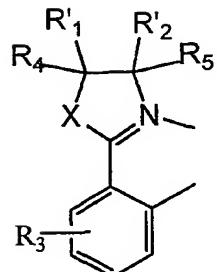
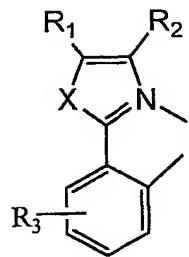
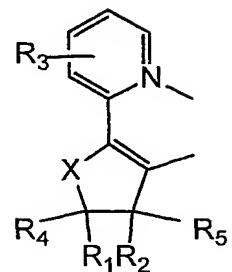
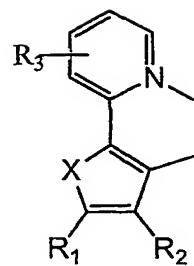
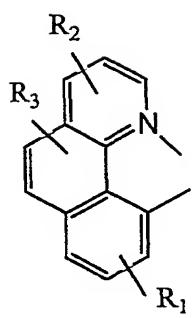
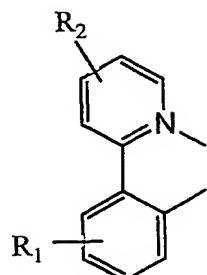
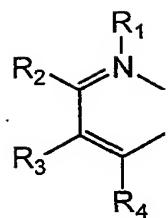


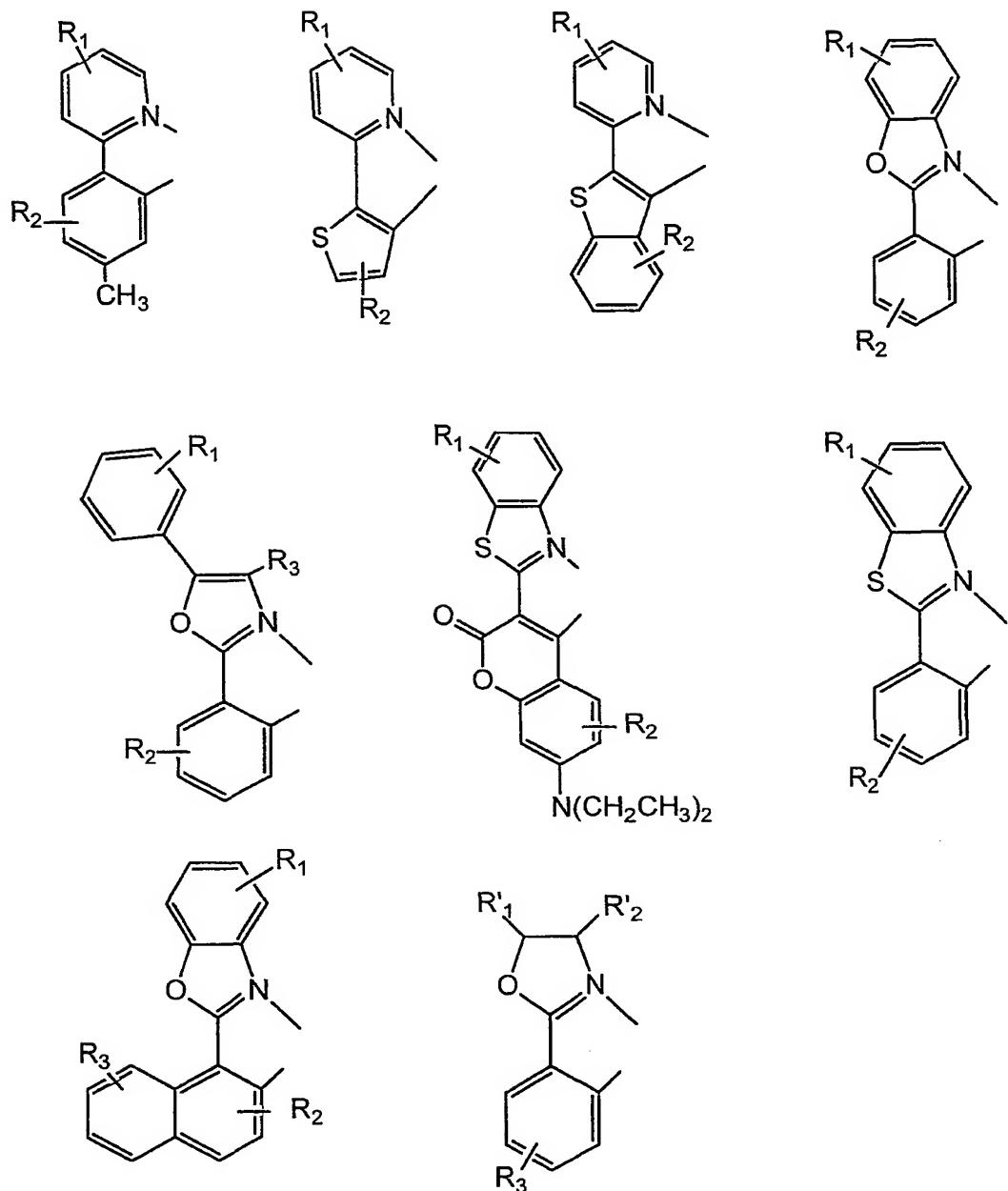
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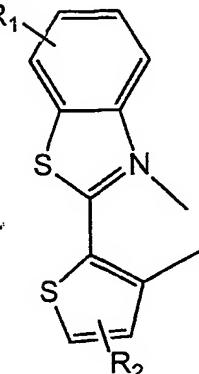
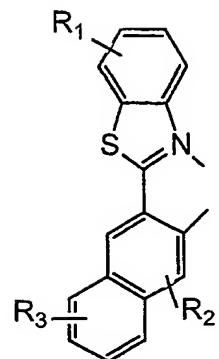
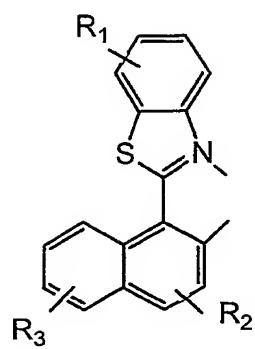
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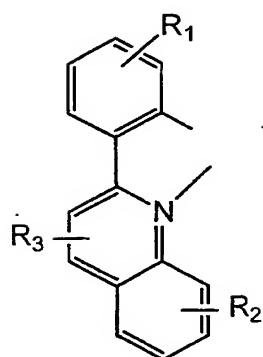
9. An organic light emitting device including an emissive layer comprising an organometallic compound comprised of:
a heavy transition metal selected from the group consisting of Os, Ir, Pt and Au;
at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the at least one mono-anionic, bidentate, carbon-coordination ligand is selected from the group consisting of :





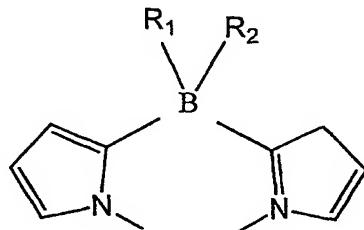
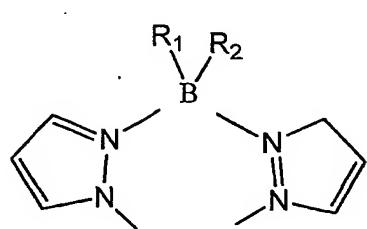
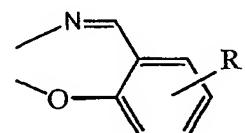
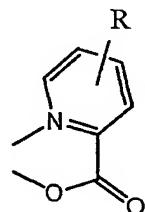
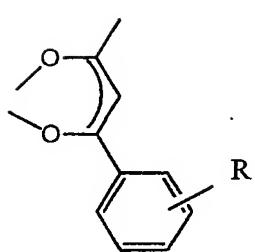
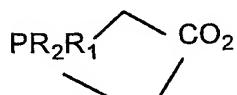
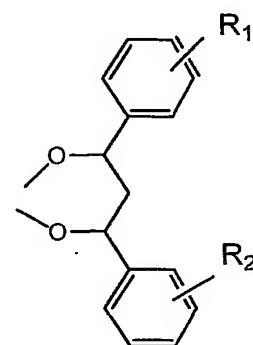
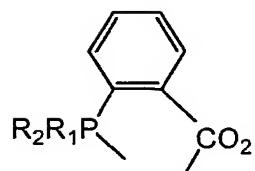
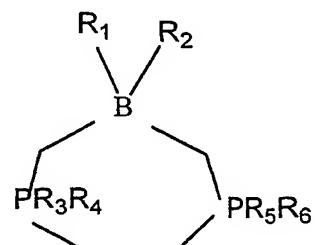
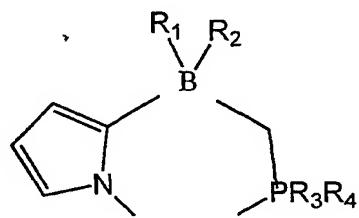


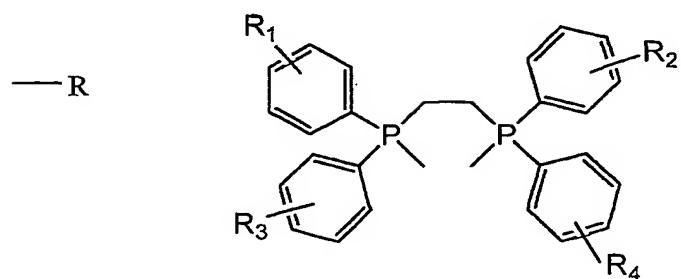
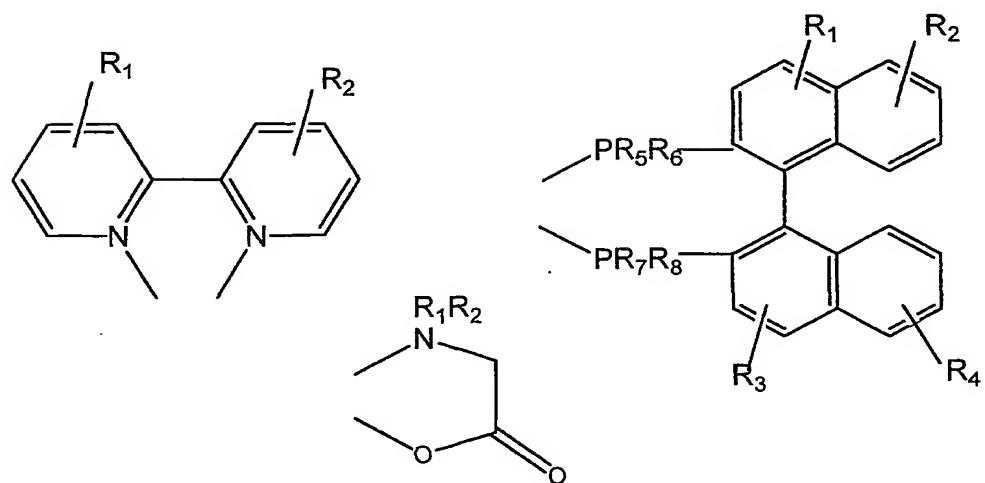
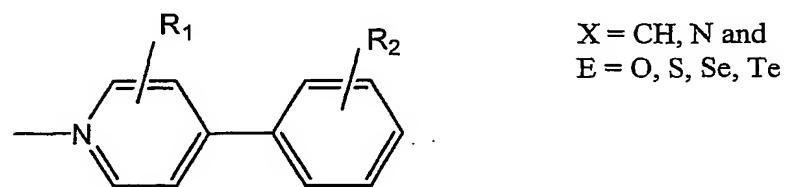
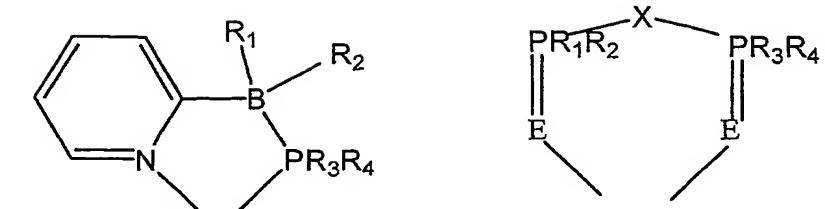
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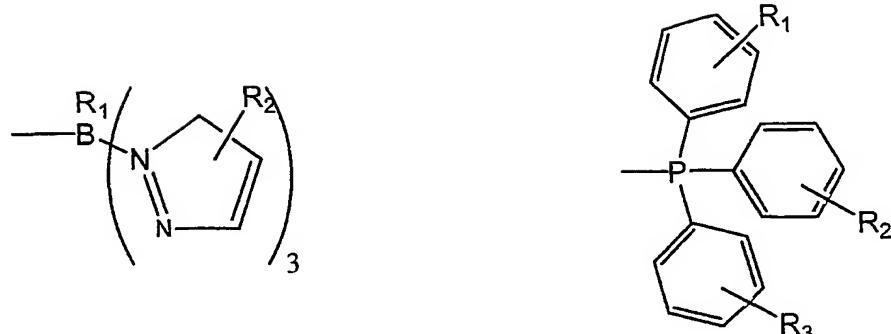


wherein X = S, O, NR; and R₁, R₂, R₃, R₄ and R₅ are, independently, hydrogen, halogen, alkyl, aryl or arylene; and R'₁ and R'₂ may, in combination, be aryl; and

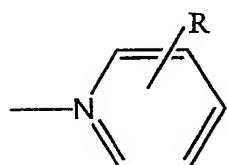
at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the at least one non-mono-anionic, bidentate, carbon-coordination ligand is selected from the group consisting of:







and



wherein R, R₁, R₂, R₃, R₄, R₅, R₆, R₇ and R₈ are, independently, hydrogen, halogen, alkyl or aryl.

10. The organic light emitting device of claim 1, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of

the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

11. The organic light emitting device of claim 2, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

12. The organic light emitting device of claim 3, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

13. The organic light emitting device of claim 4, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

14. The organic light emitting device of claim 5, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

15. The organic light emitting device of claim 6, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

16. The organic light emitting device of claim 7, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

17. The organic light emitting device of claim 8, wherein the emissive layer further comprises:

a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.

18. The organic light emitting device of claim 9, wherein the emissive layer further comprises:
a host material having a lowest triplet excited state having a first decay rate of less than about 1 per second; wherein the organometallic compound is present as a guest material dispersed in the host material, the organometallic compound having a lowest triplet excited state having a radiative decay rate of greater than about 1×10^5 per second and wherein the energy level of the lowest triplet excited state of the host material is lower than the energy level of the lowest triplet excited state of the organometallic compound.
19. The organic light emitting device of claim 10, wherein the energy difference between the lowest triplet excited state of the organometallic compound and a

corresponding relaxed stated of the organometallic compound has a corresponding wavelength of about 420 nm to 480 nm for blue light emission.

20. The organic light emitting device of claim 10, wherein the energy difference between the lowest triplet excited state of the organometallic compound and a corresponding relaxed stated of the organometallic compound has a corresponding wavelength of about 480 nm to 510 nm for aqua-blue light emission.
21. The organic light emitting device of claim 10, wherein the host material has a bandgap with an energy difference corresponding to about 470 nm and the organometallic compound has a lowest triplet excited state at an energy level at about 450 nm.
22. The organic light emitting device of claim 10, wherein the host material is an electron transport layer.
23. The organic light emitting device of claim 10, wherein the host material conducts electrons primarily through hole transmission.
24. The organic light emitting device of claim 10, wherein the ratio of the host material and organometallic compound decay rates is at least about 1:1000 to about 5:1000.
25. The organic light emitting device of claim 10, wherein the host material is TPD.

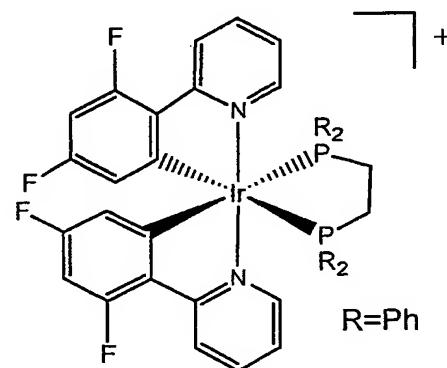
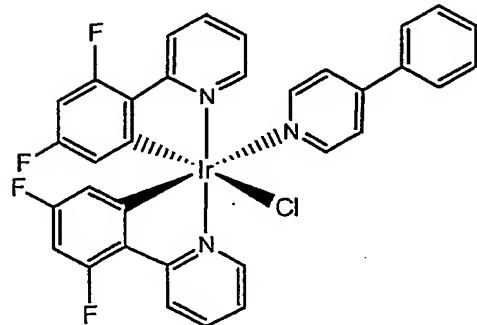
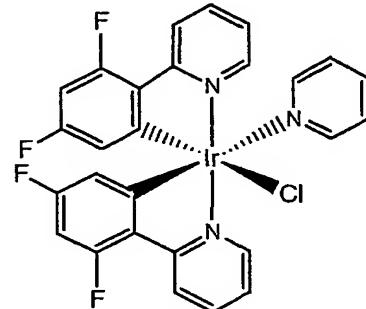
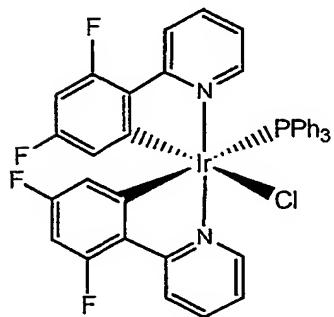
26. The organic light emitting device of claim 10, wherein a plurality of organometallic compounds are dispersed in the host material.
27. An organometallic compound comprising:
a heavy transition metal that produces an efficient phosphorescent emission at room temperature from a mixture of metal-to-ligand charge transfer and $\pi - \pi^*$ ligand states;
at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the at least one mono-anionic, bidentate, carbon-coordination ligand is substituted with at least one of an electron donating substituent and an electron withdrawing substituent, wherein the at least one of an electron donating substituent and an electron withdrawing substituent shifts the emission, relative to an un-substituted mono-anionic, bidentate, carbon-coordination ligand, to either the blue, green or red region of the visible spectrum;
and
at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the at least one non-mono-anionic, bidentate, carbon-coordination ligand causes the emission to have a well defined vibronic structure.
28. The organometallic compound of claim 27, wherein the heavy transition metal is selected from the group consisting of Os, Ir, Pt and Au.

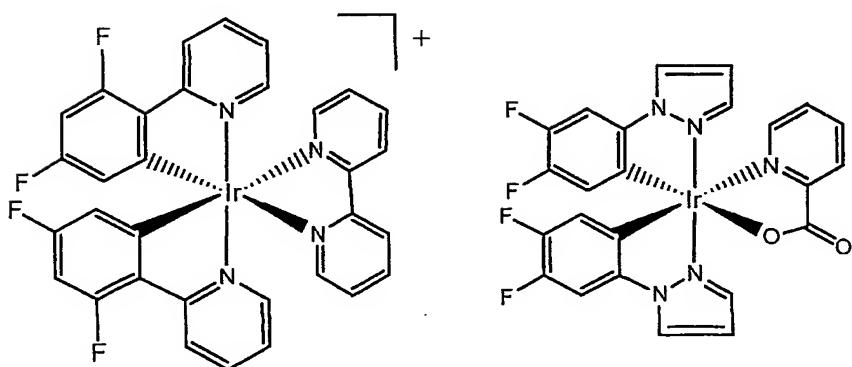
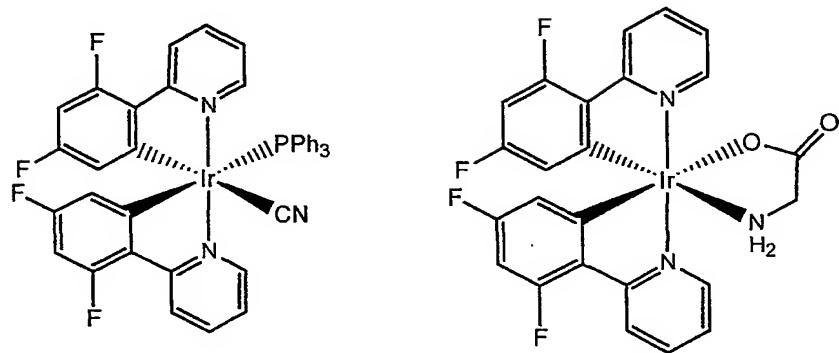
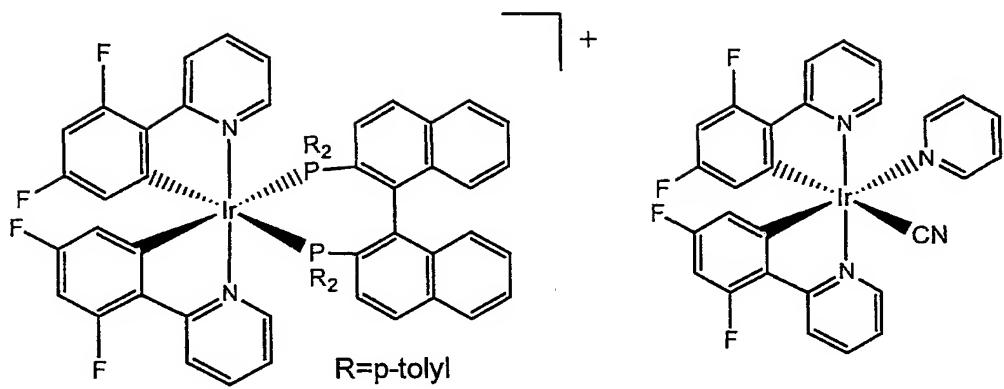
29. An organometallic compound comprising:

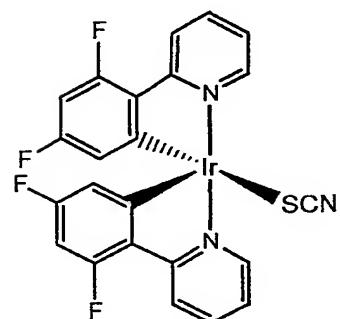
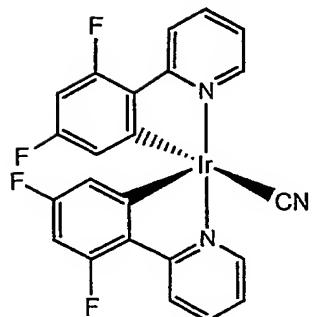
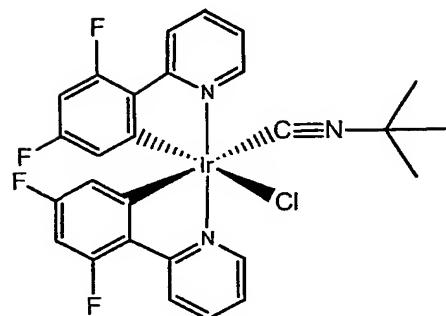
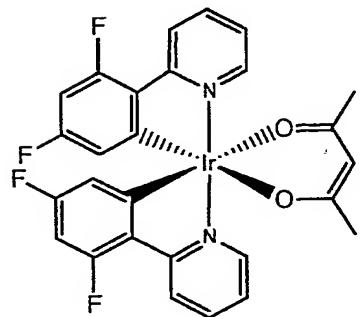
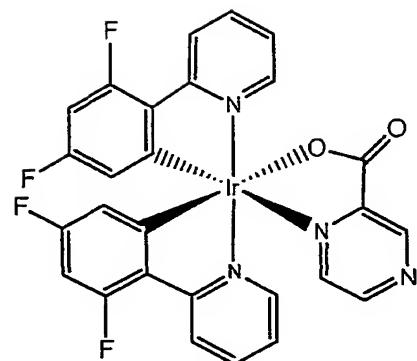
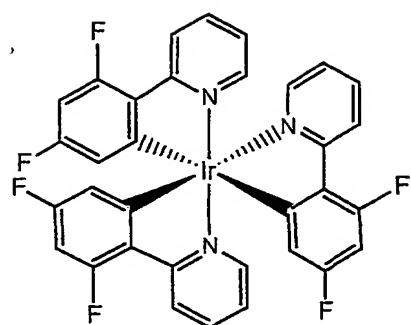
a heavy transition metal;

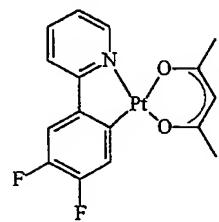
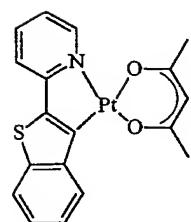
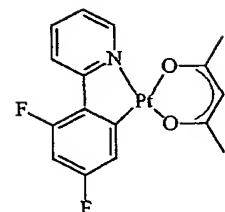
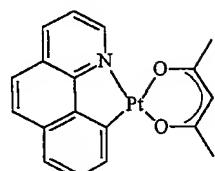
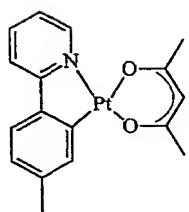
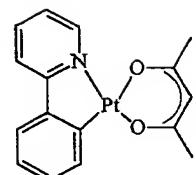
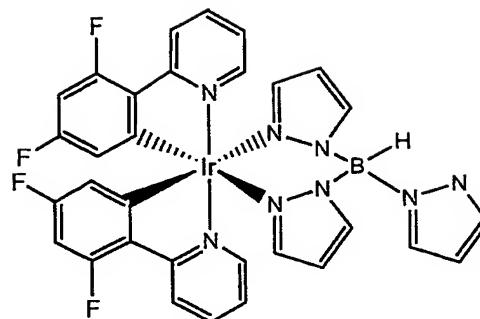
at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal; and

at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the organometallic compound has a chemical structure represented by a formula selected from the group consisting of:

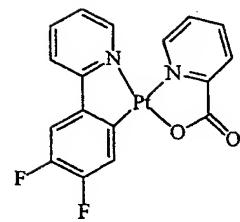








and

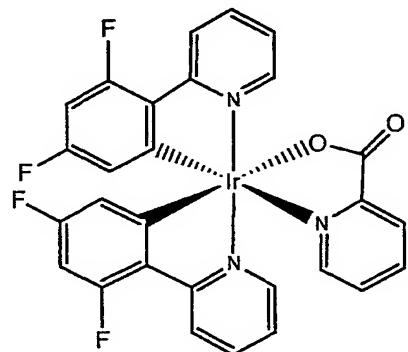


30. An organometallic compound comprising:

a heavy transition metal;

at least one mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal; and

at least one non-mono-anionic, bidentate, carbon-coordination ligand bound to the heavy transition metal, wherein the organometallic compound has a chemical structure represented by the following formula:



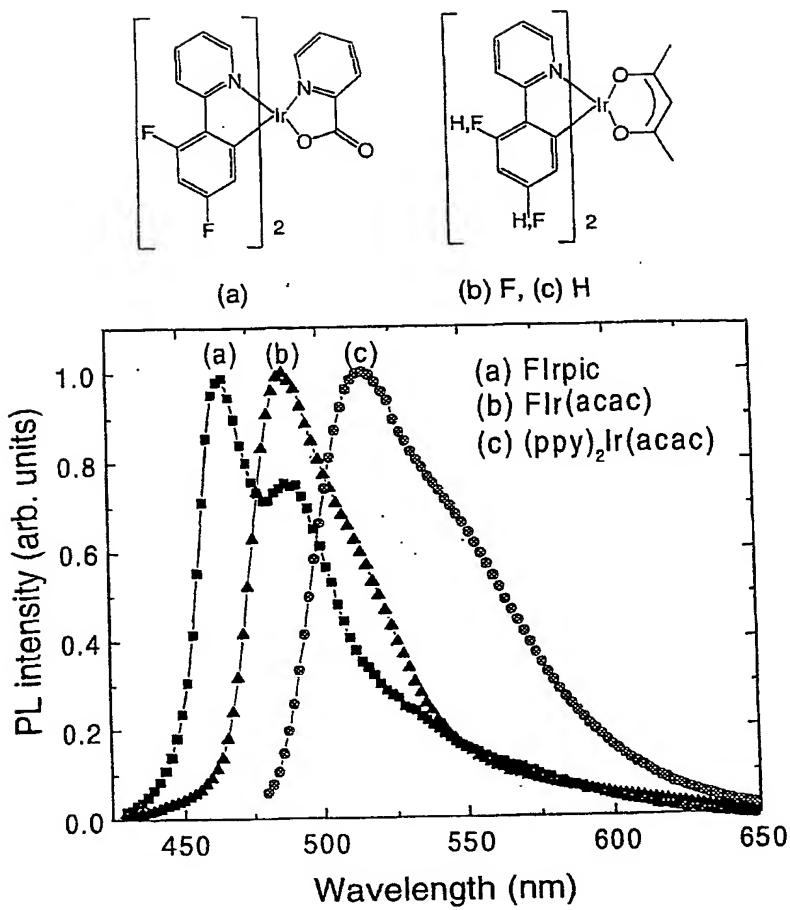


Figure 1a:

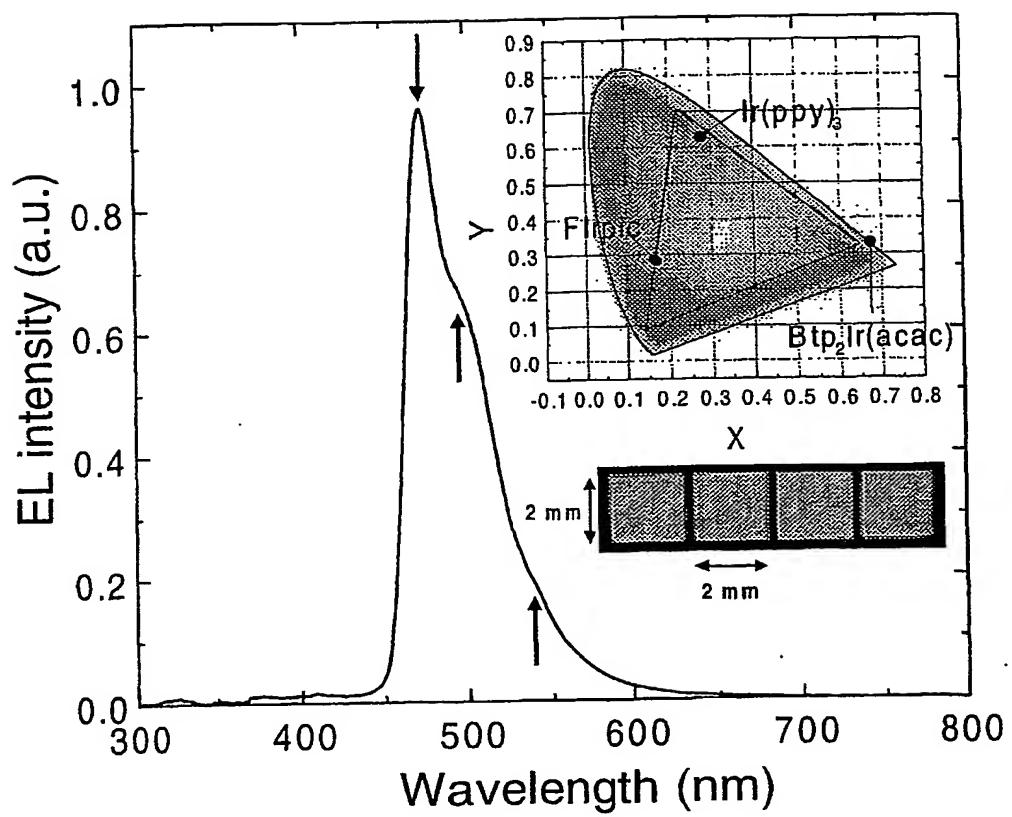


Figure 1b:

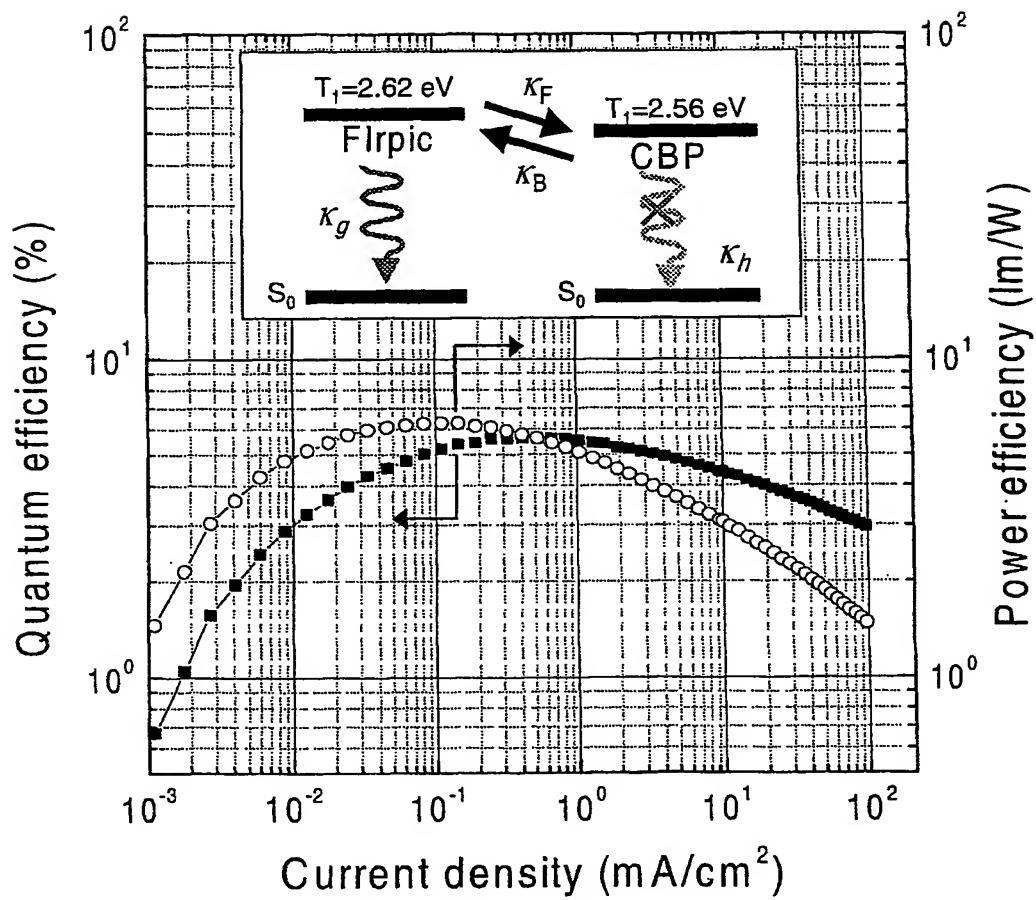


Figure 2:

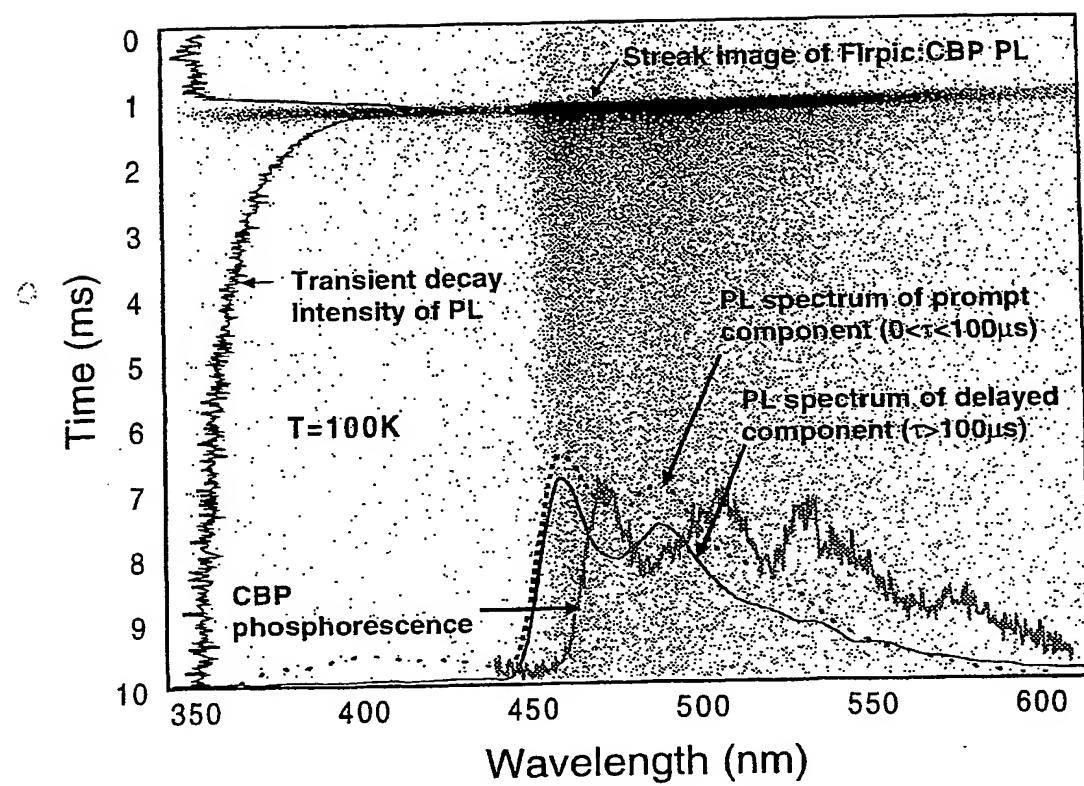


Figure 3:

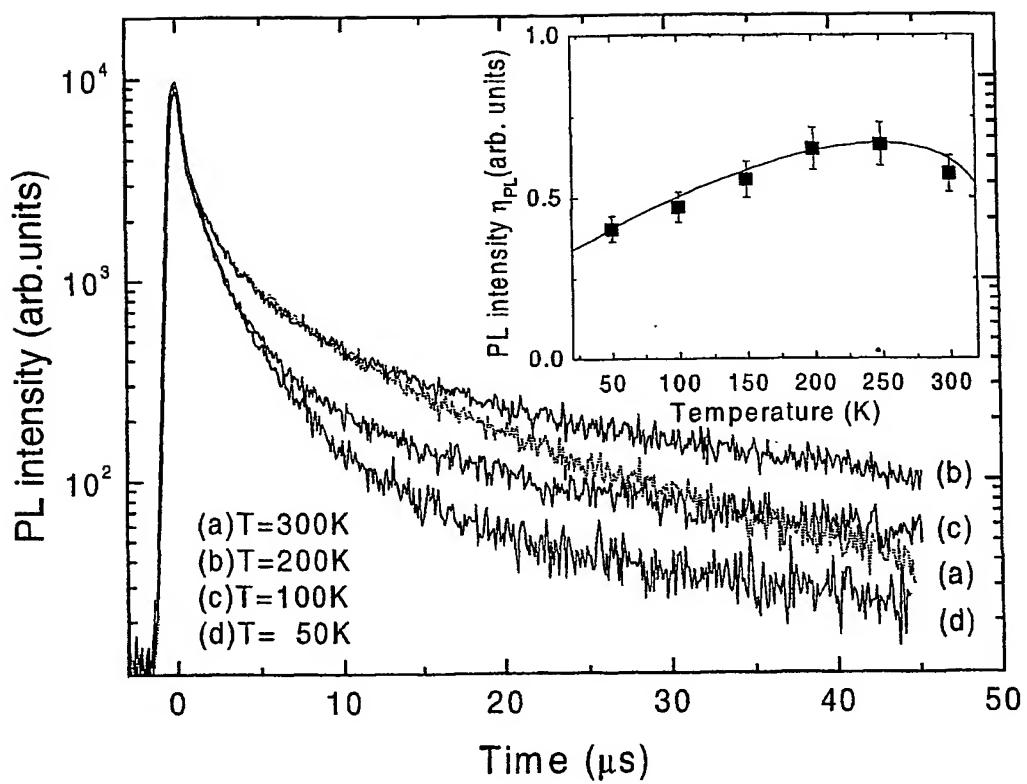
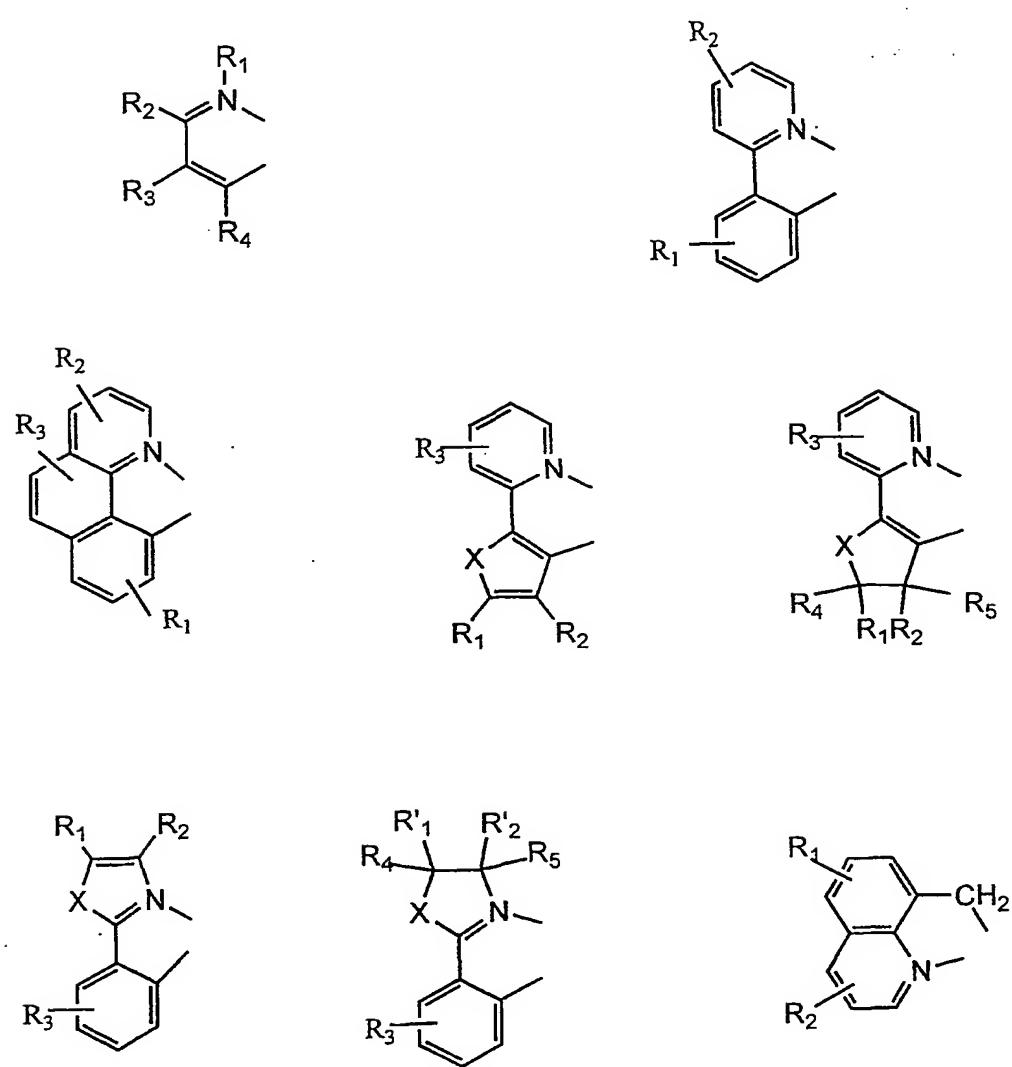


Figure 4:

figure 5a

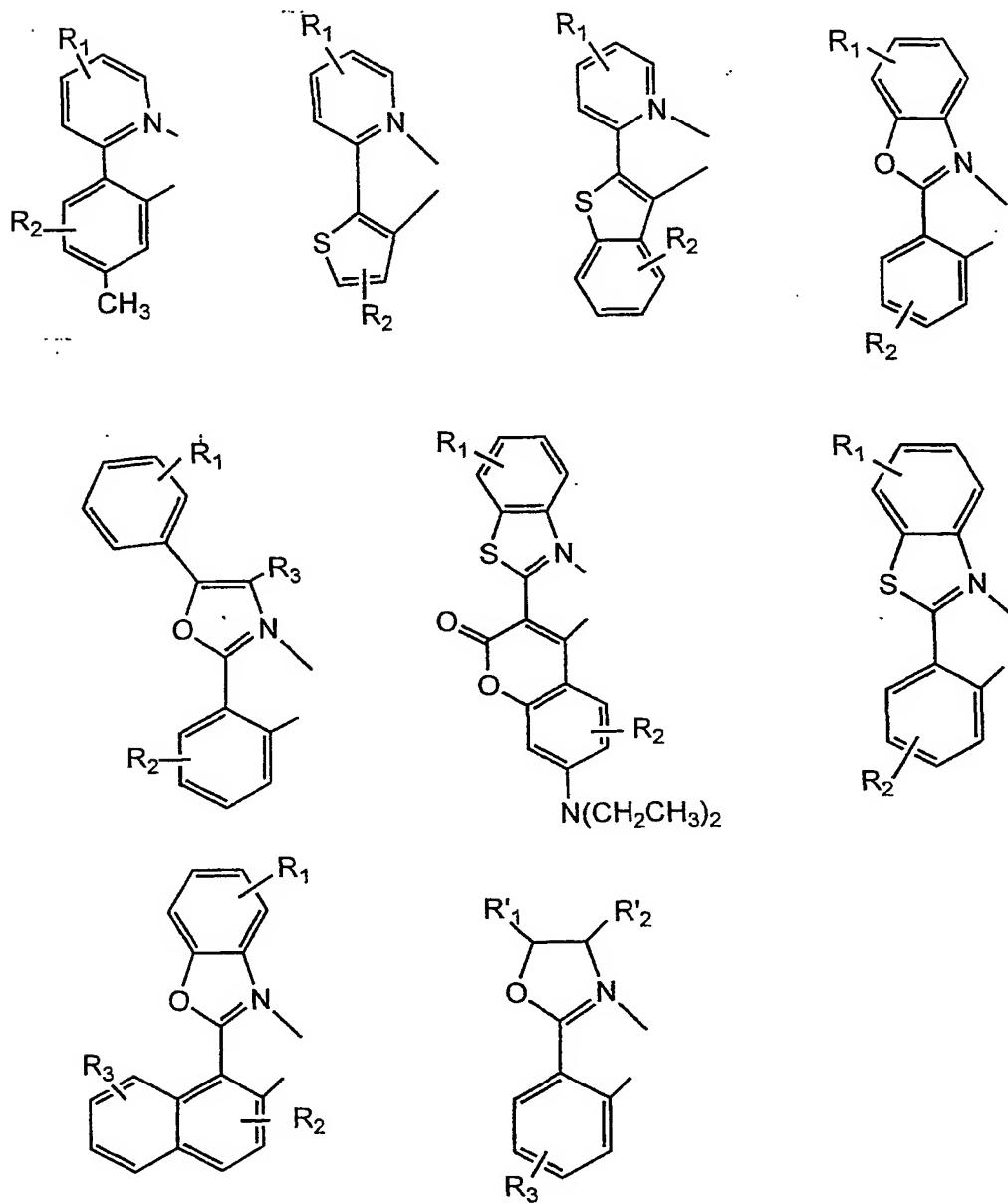
Generic Mono-Anionic, Bidentate, Carbon-Coordination Ligands-I



X = S, O, NR; and R₁, R₂, R₃, R₄ and R₅ are, independently, hydrogen, halogen, alkyl, aryl or arylene; and R'₁ and R'₂ may, in combination, be aryl.

Figure 5b
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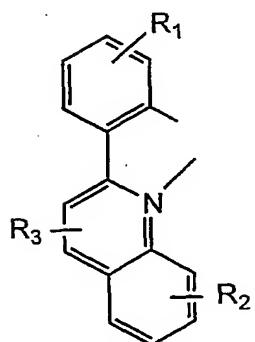
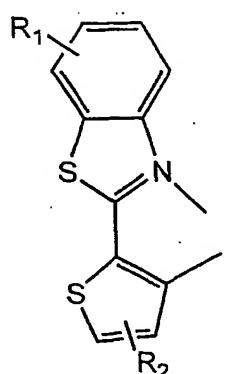
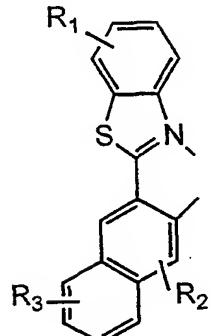
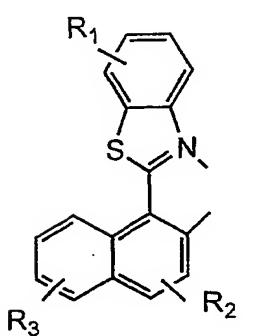
Generic Mono-Anionic, Bidentate, Carbon-Coordination Ligands-II



X = S, O, NR; and R₁, R₂, R₃, R₄ and R₅ are, independently, hydrogen, halogen, alkyl, aryl or arylene; and R'₁ and R'₂ may, in combination, be aryl.

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Figure 5c

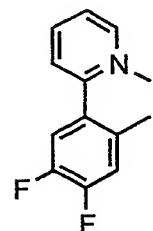
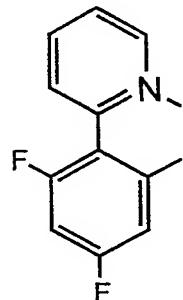
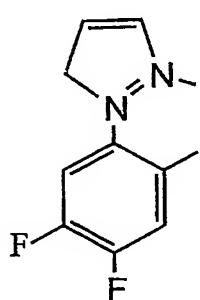
Generic Mono-Anionic, Bidentate, Carbon-Coordination Ligands-III



X = S, O, NR; and R₁, R₂, R₃, R₄ and R₅ are, independently, hydrogen, halogen, alkyl, aryl or arylene.

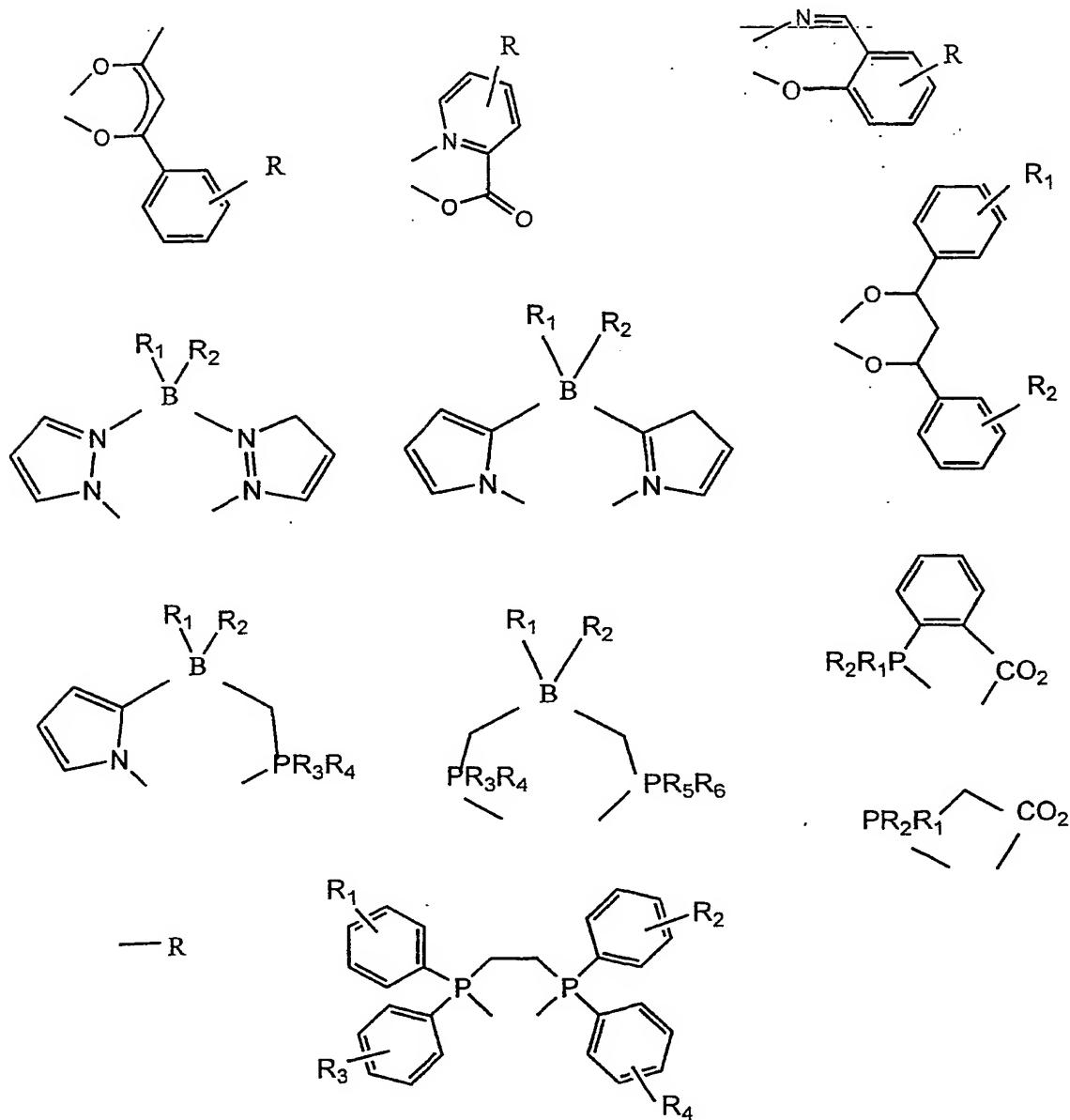
figure 5d

Specific Mono-Anionic, Bidentate, Carbon-Coordination Ligands-I



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Figure 6a

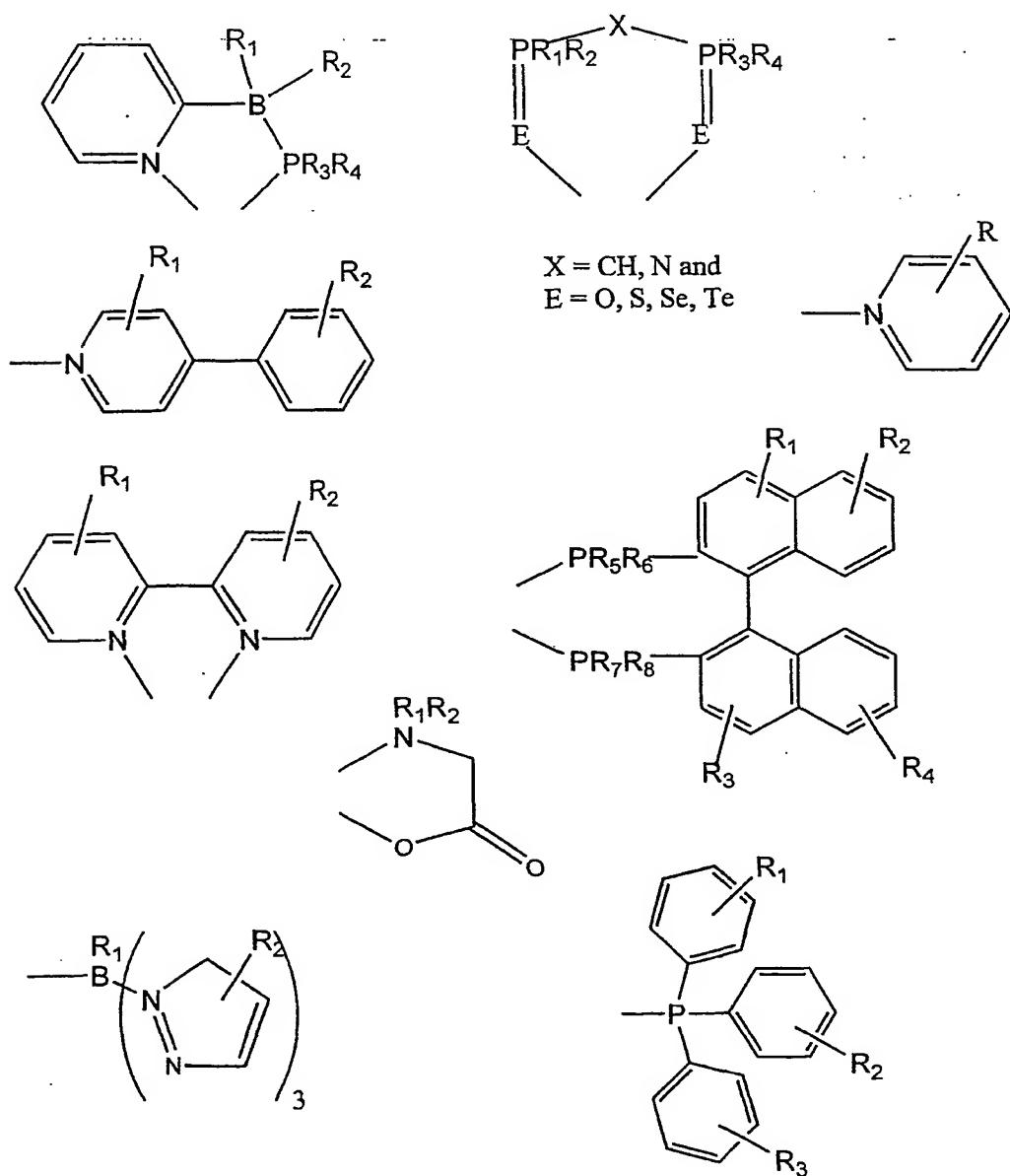
Generic Non-Mono-Anionic, Bidentate, Carbon-Coordination Ligands-I



R, R₁, R₂, R₃, R₄, R₅, and R₆ are, independently, hydrogen, halogen, alkyl or aryl.

Figure 6b

Generic Non-Mono-Anionic, Bidentate, Carbon-Coordination Ligands-II



R, R₁, R₂, R₃, R₄, R₅, R₆, R₇ and R₈ are, independently, hydrogen, halogen, alkyl or aryl.

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Figure 6c

Specific Non-Mono-Anionic, Bidentate, Carbon-Coordination Ligands

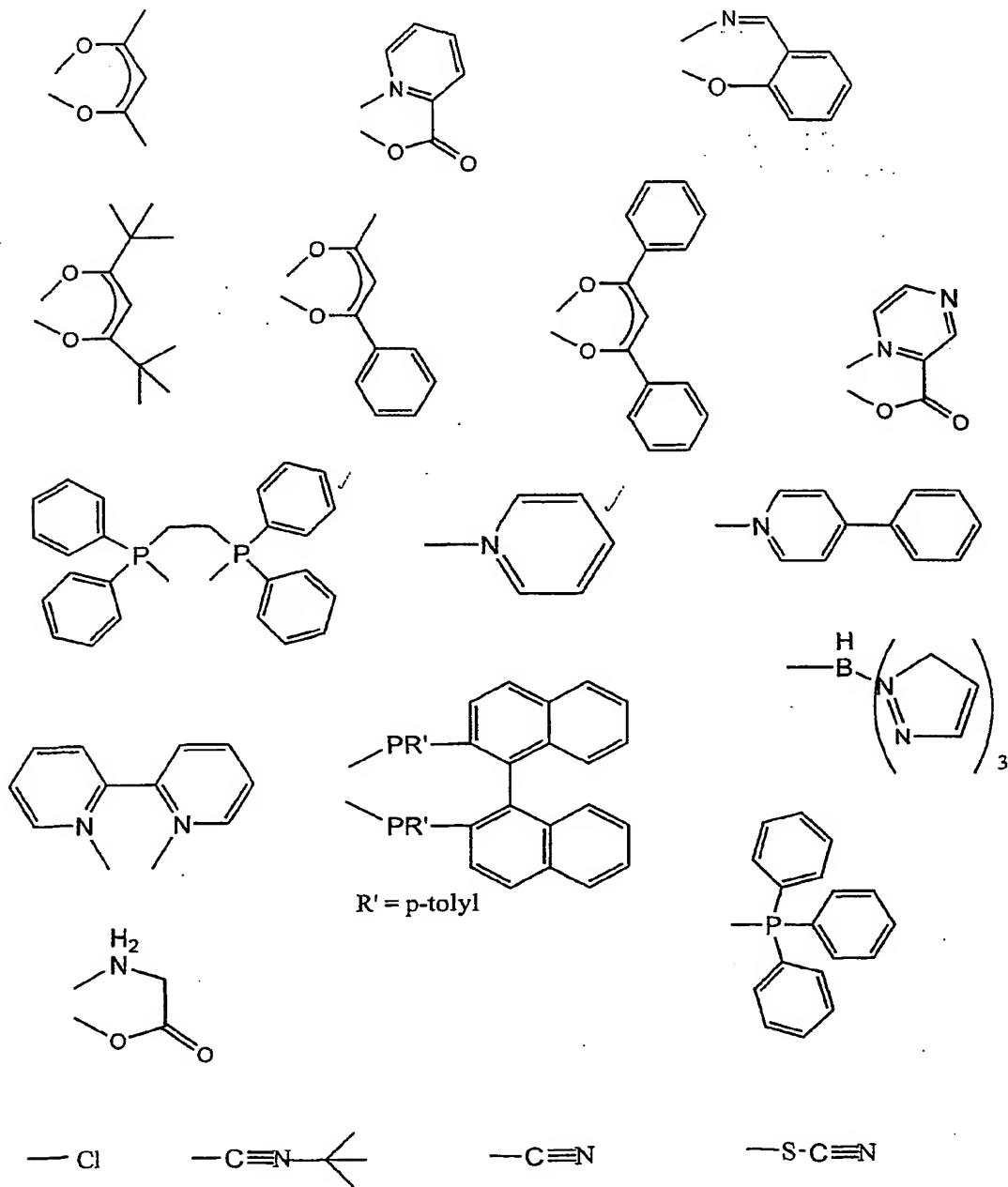


Figure 7a

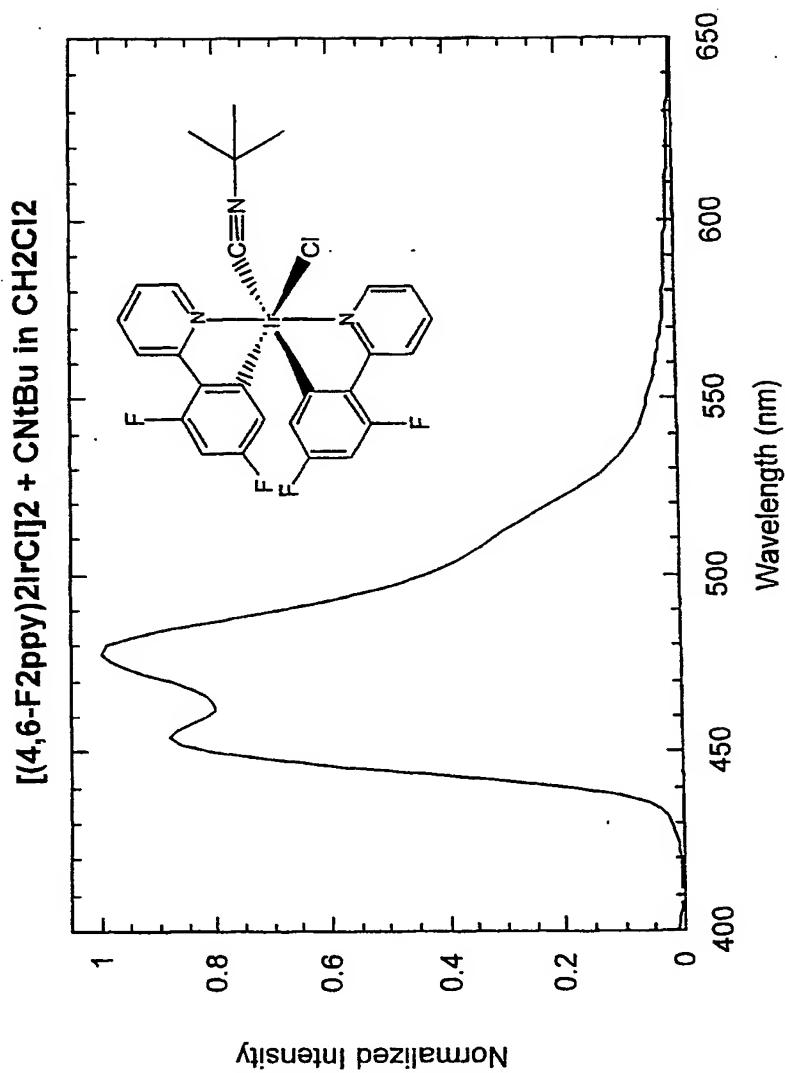


Figure 7b

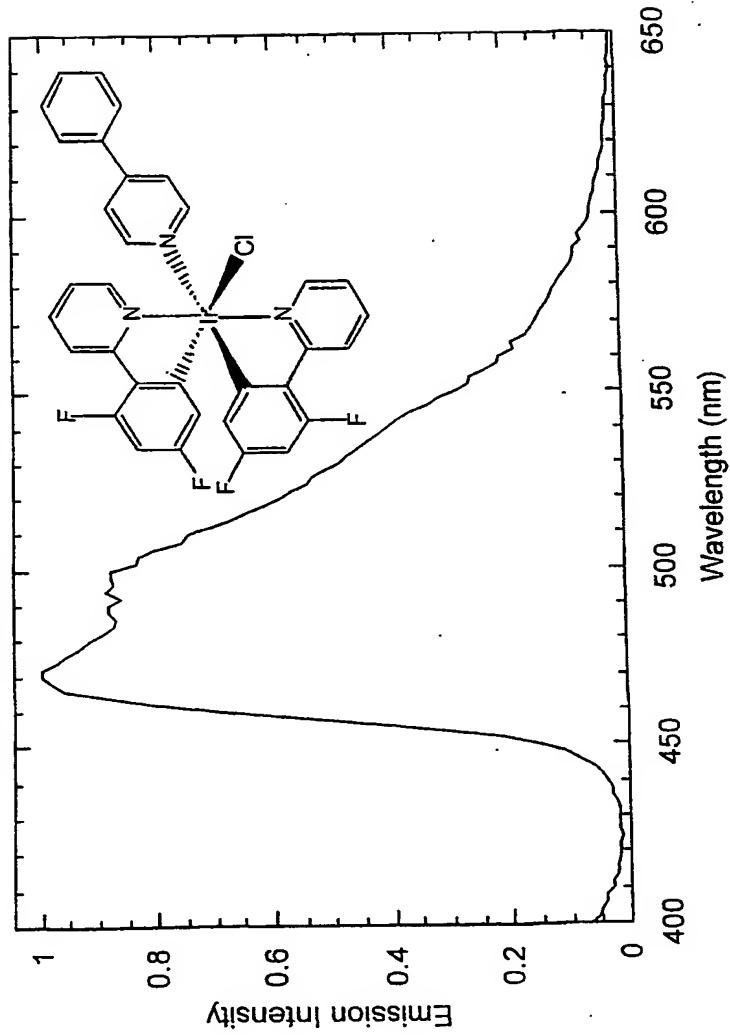
 $[\text{Ir}(4,6\text{-F}_2\text{ppy})_2\text{Cl}] + 4\text{-Phpyr}$ in CH_2Cl_2 

Figure 7c

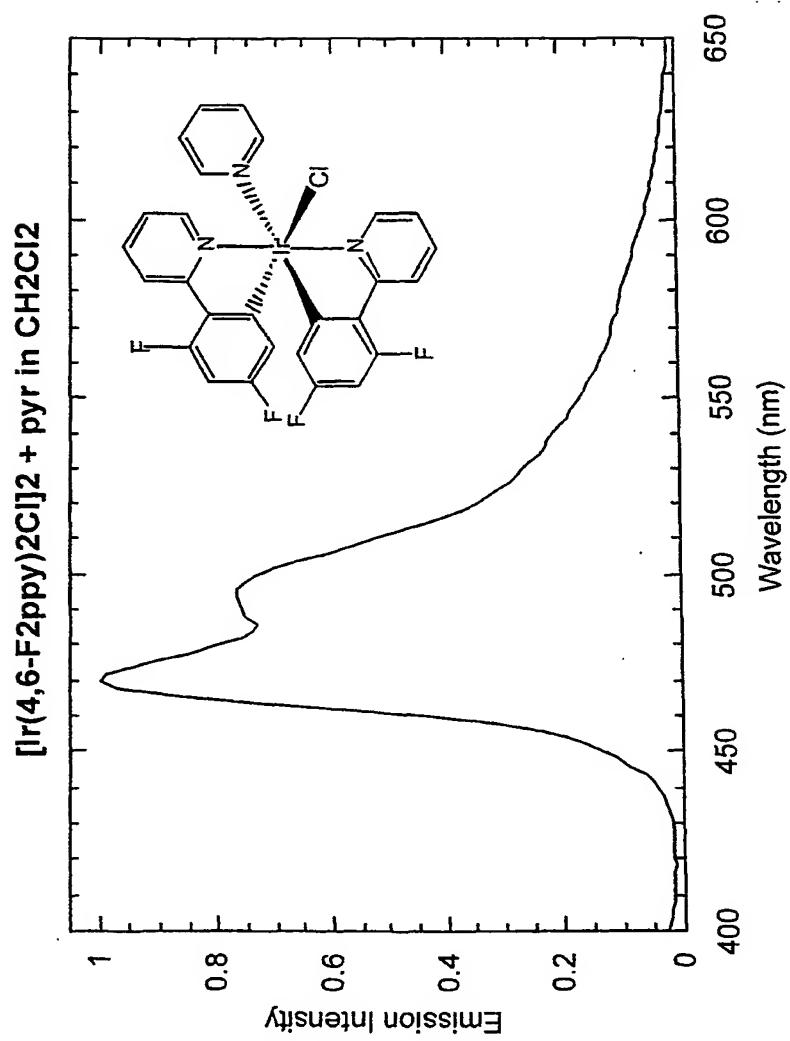


Figure 7d

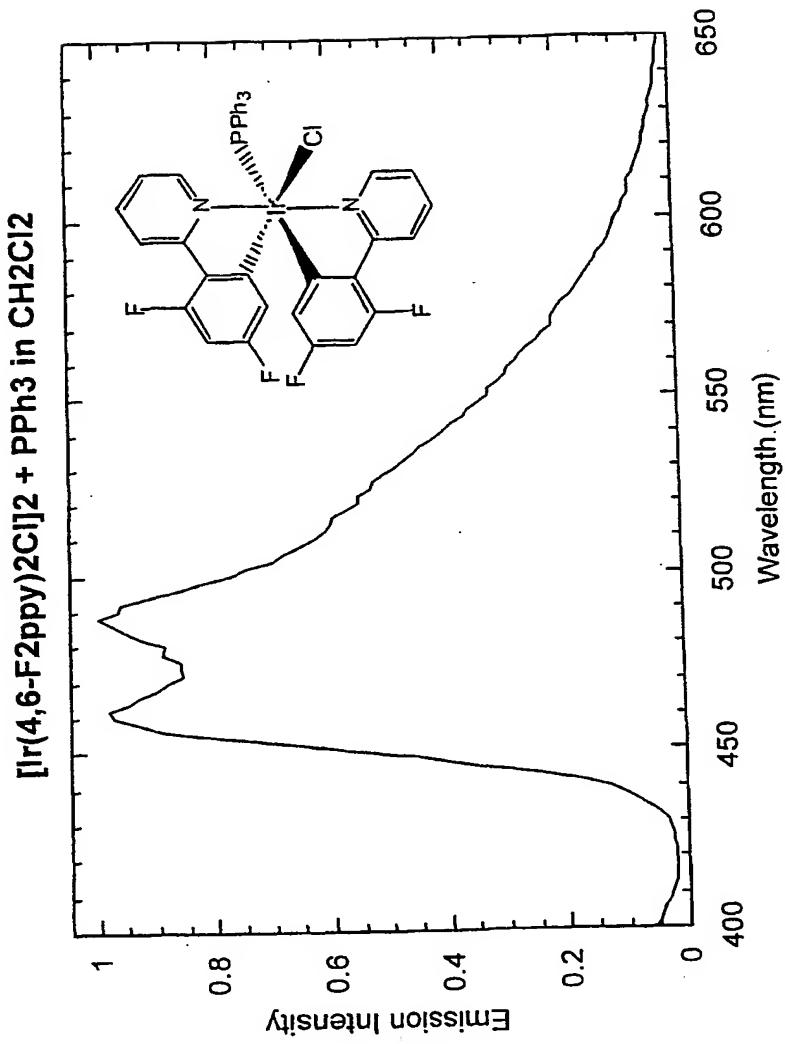


Figure 7e

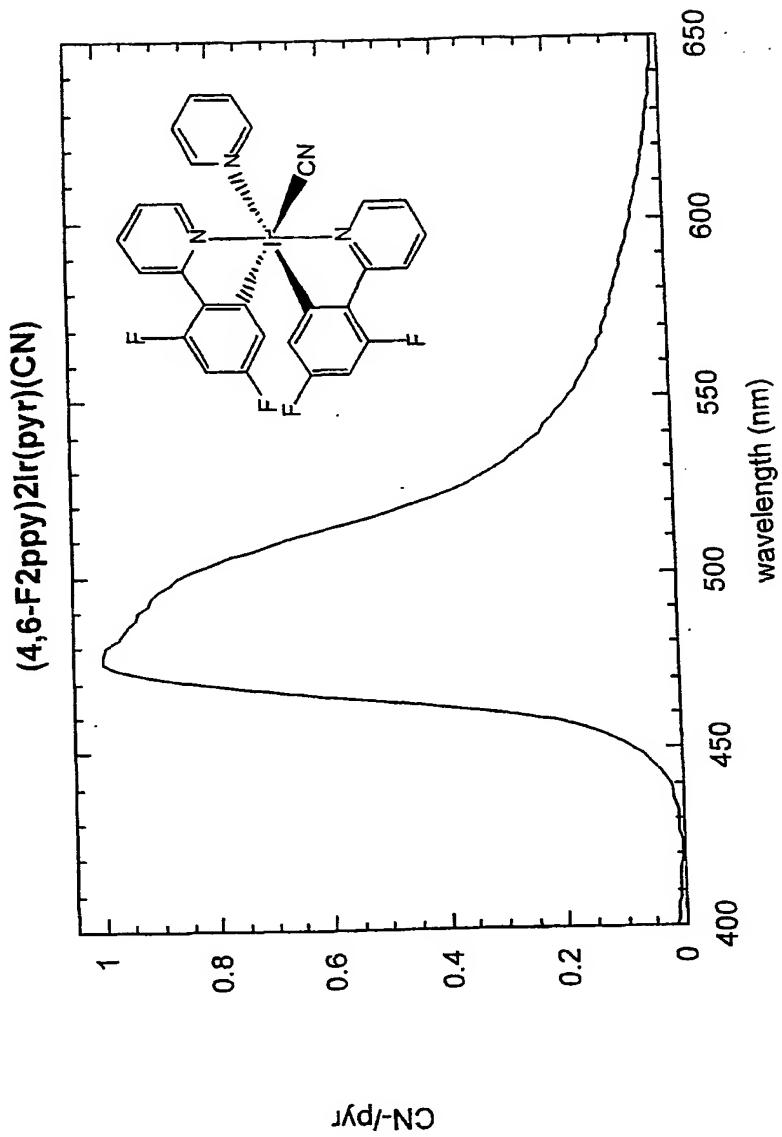


figure 7f

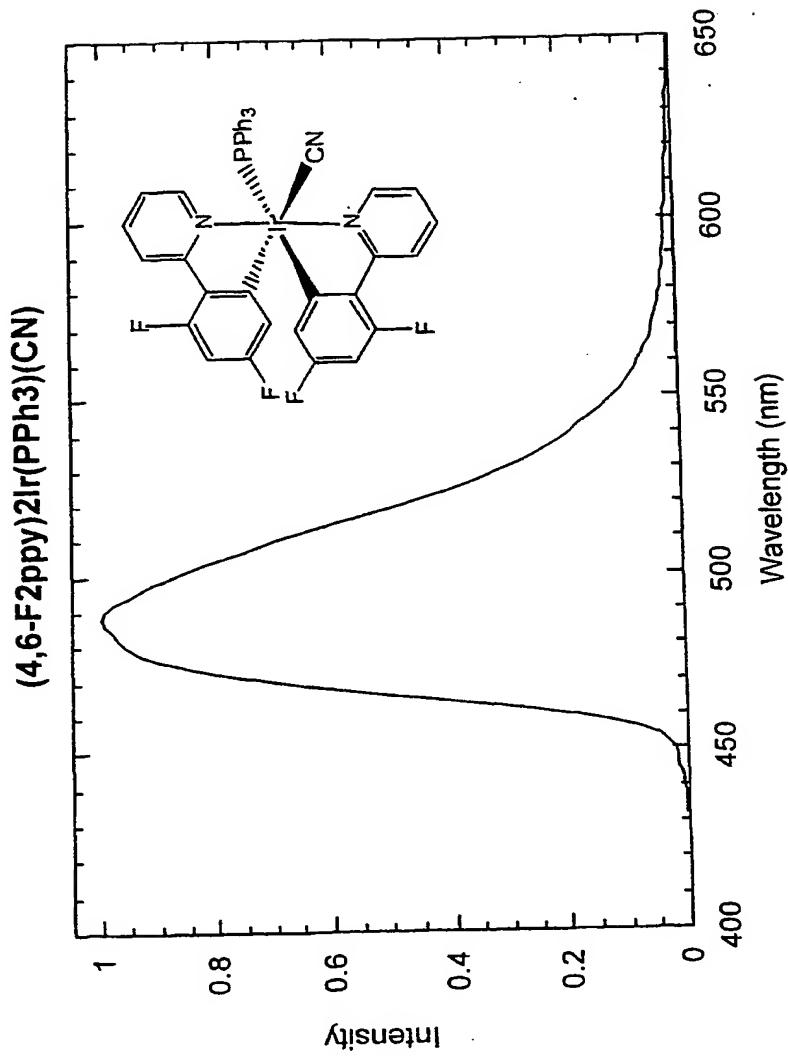


Figure 7g

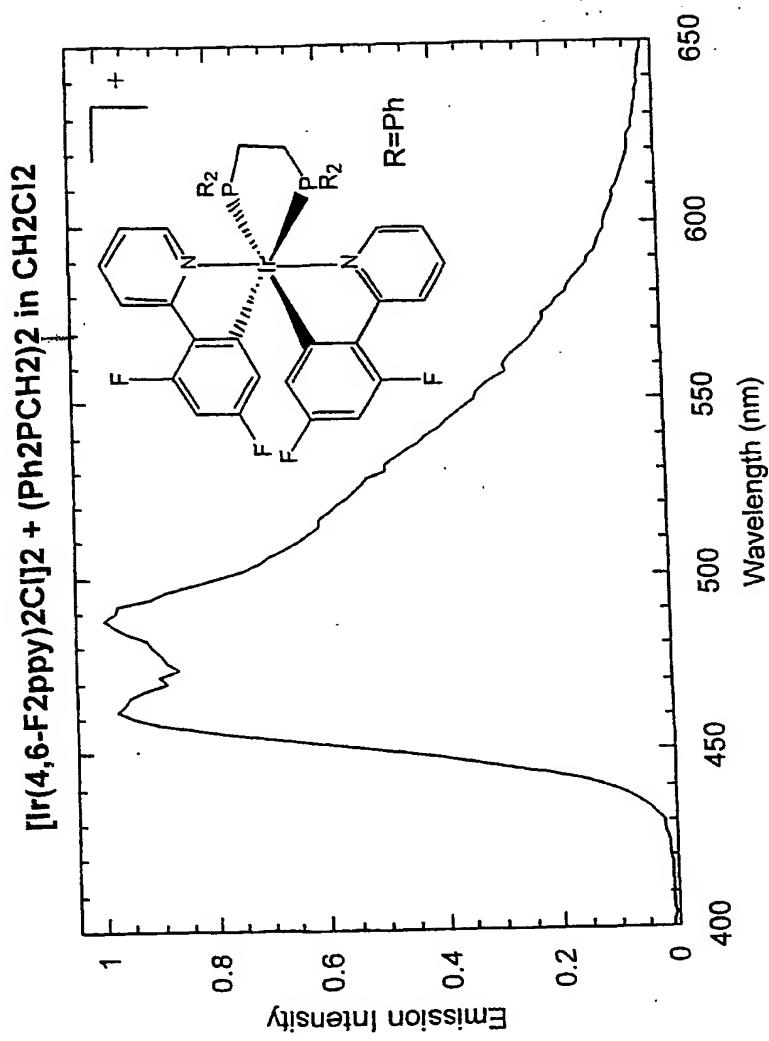


Figure 7h

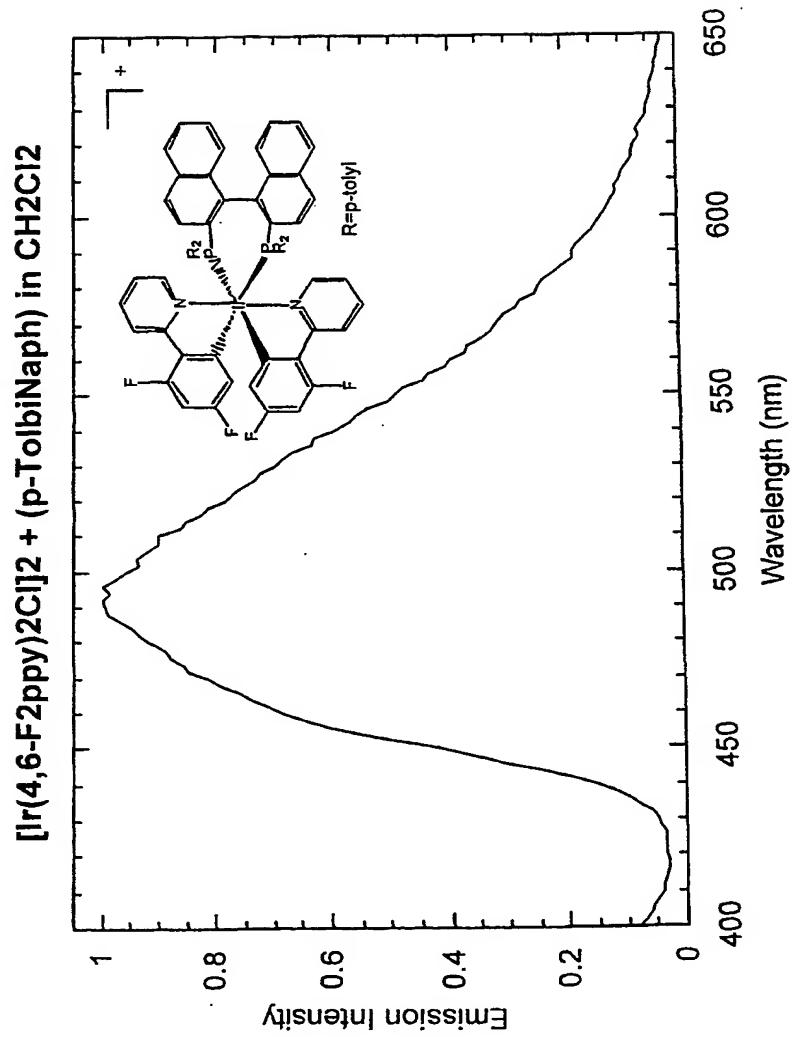


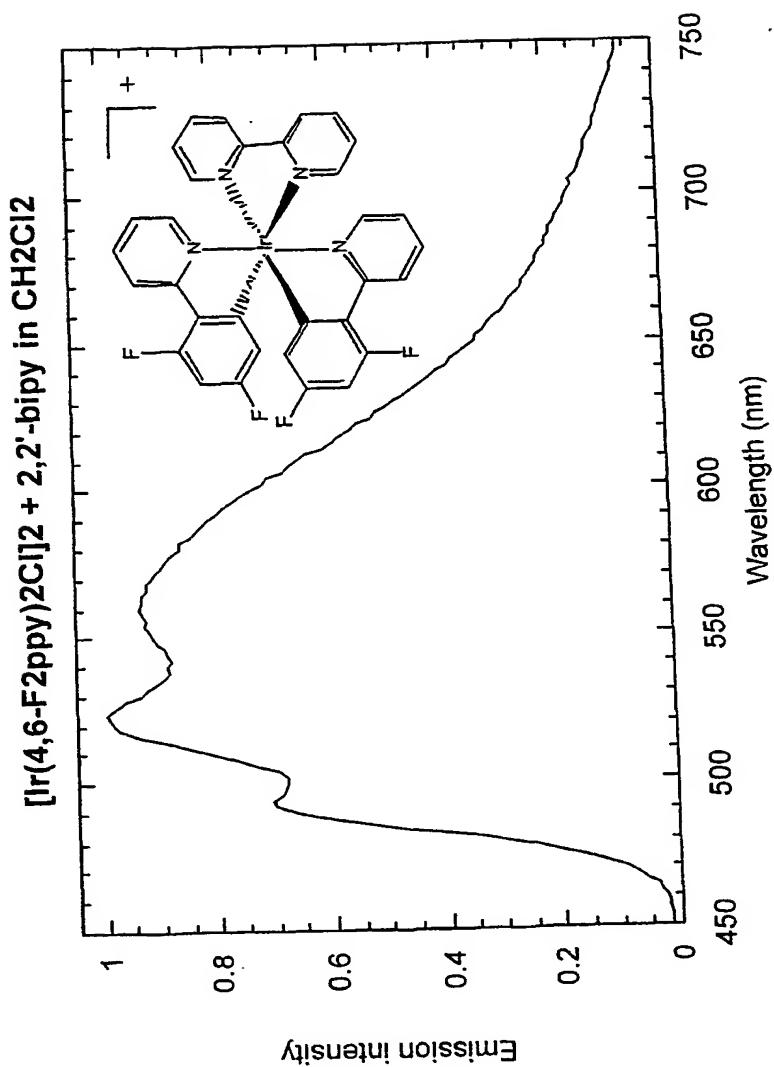
Figure 7*i*

Figure 7:

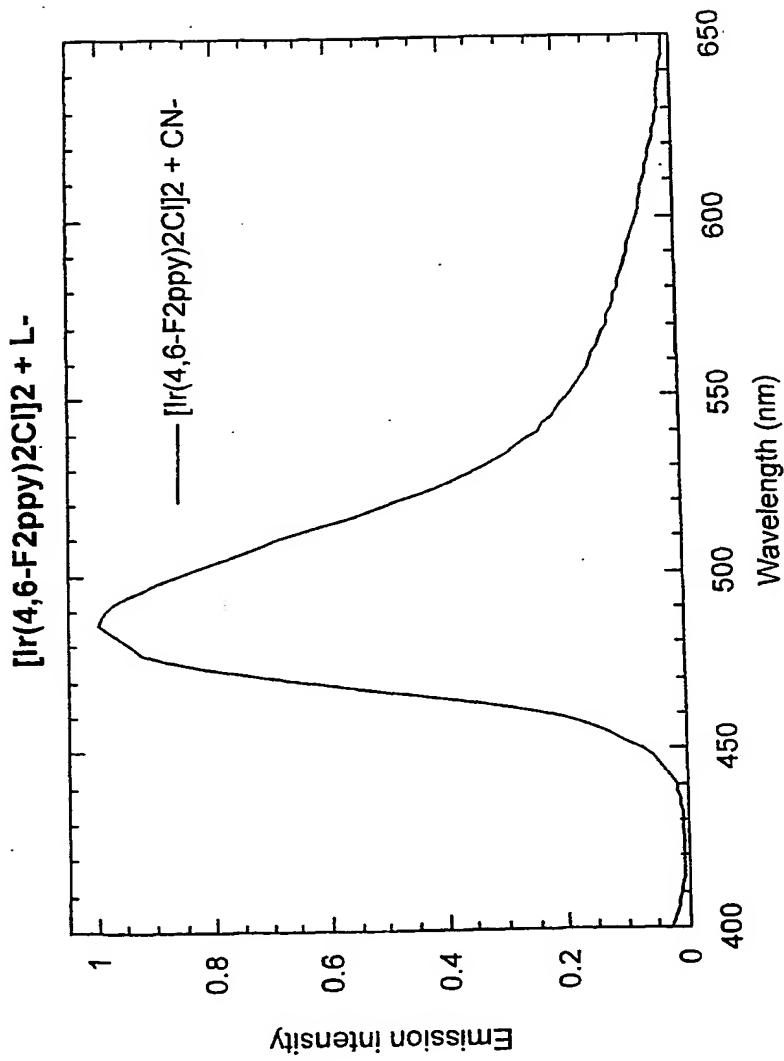


Figure 7k

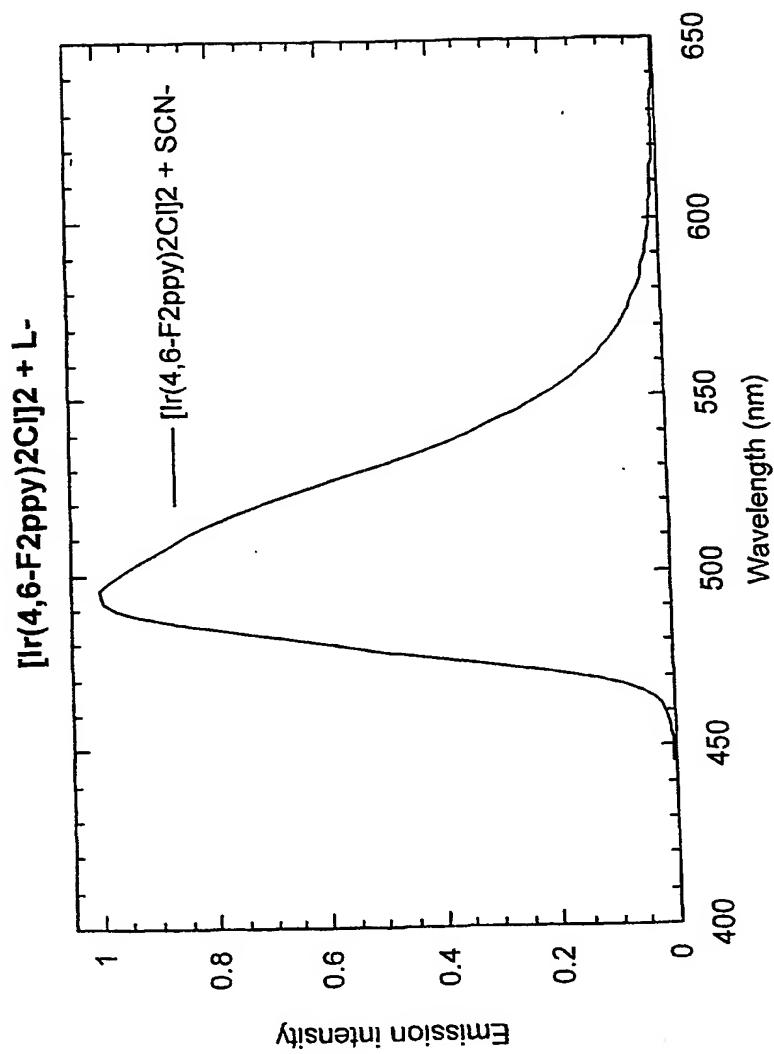


Figure 70

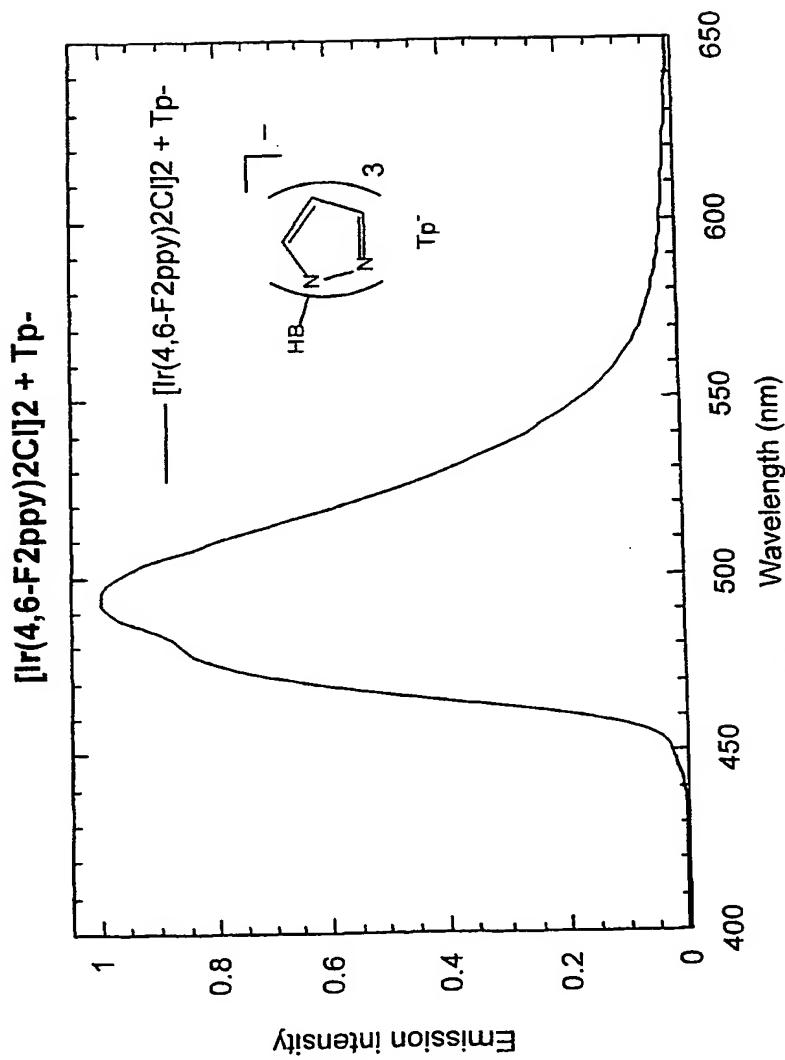


Figure 7m

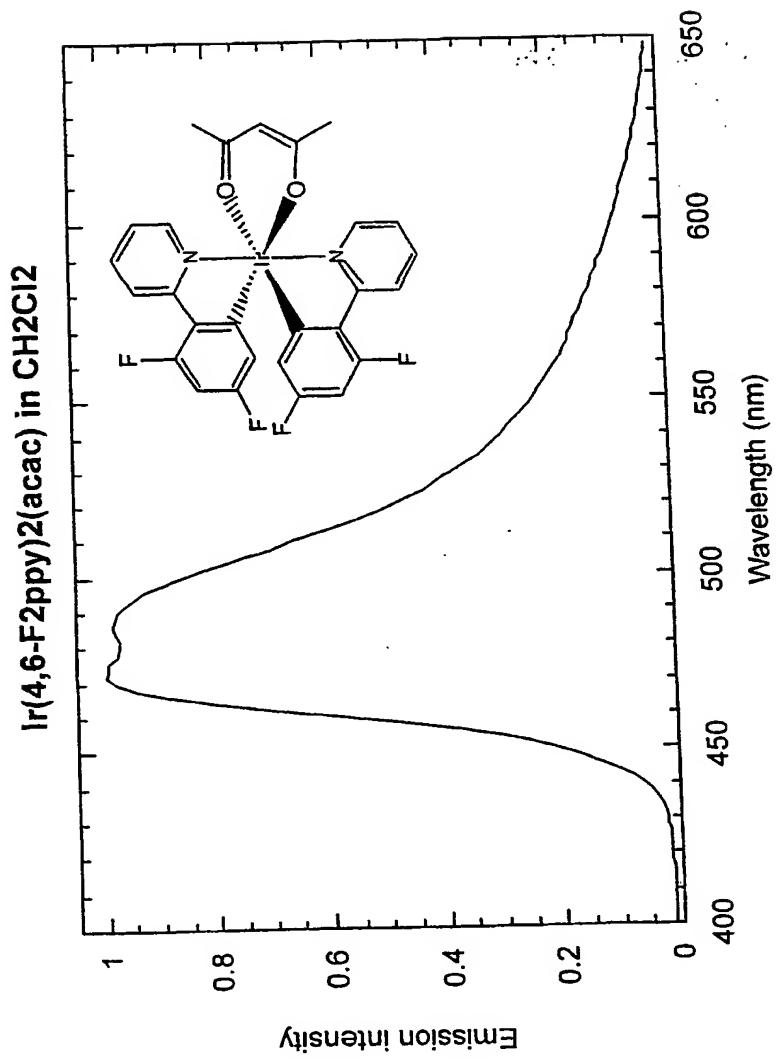


Figure 7n

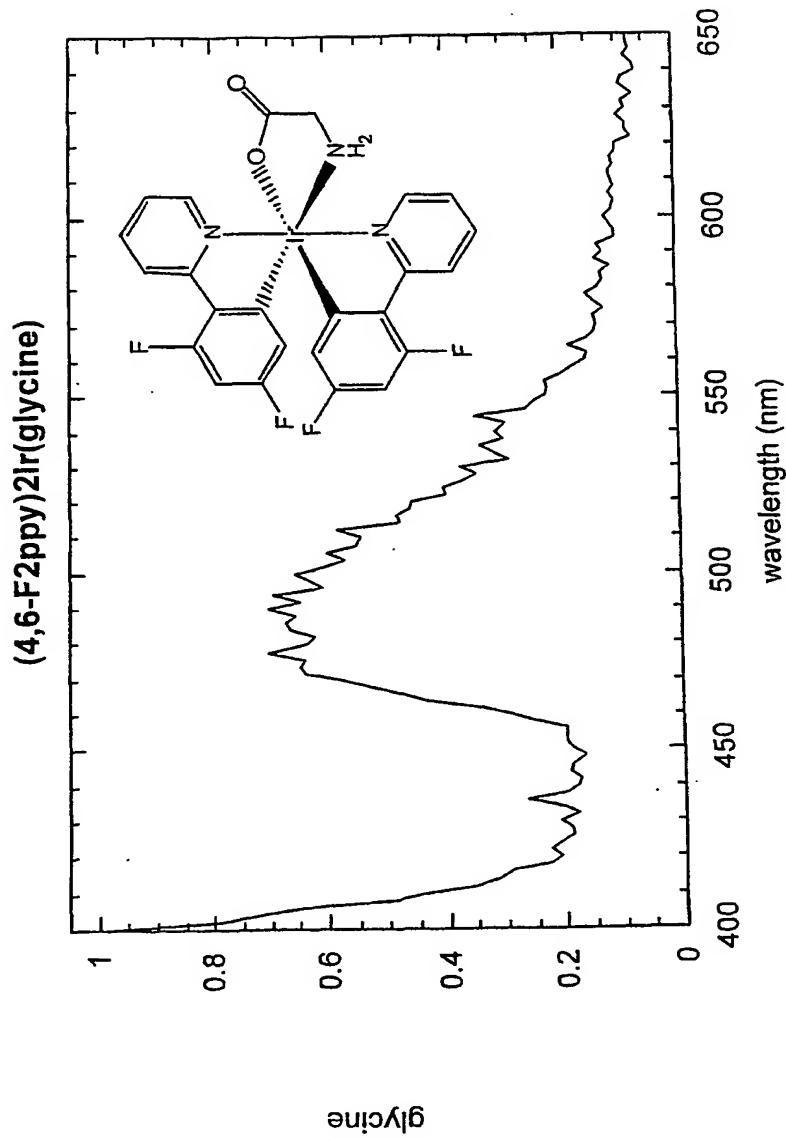


Figure 7.

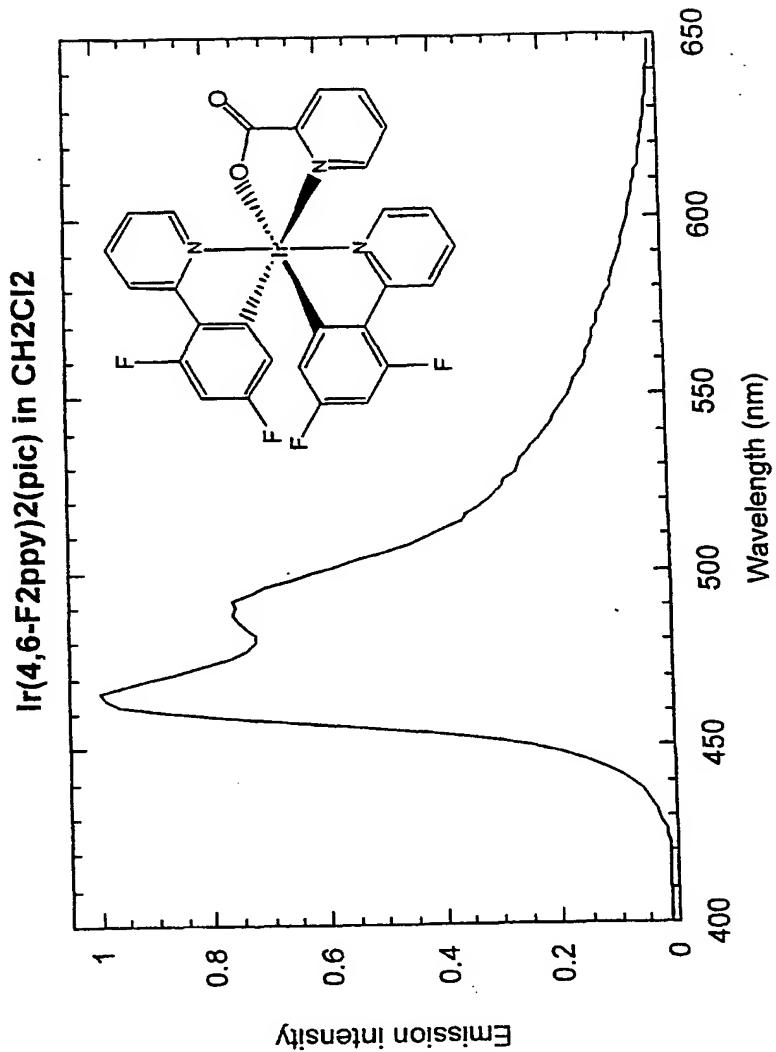


figure 7p

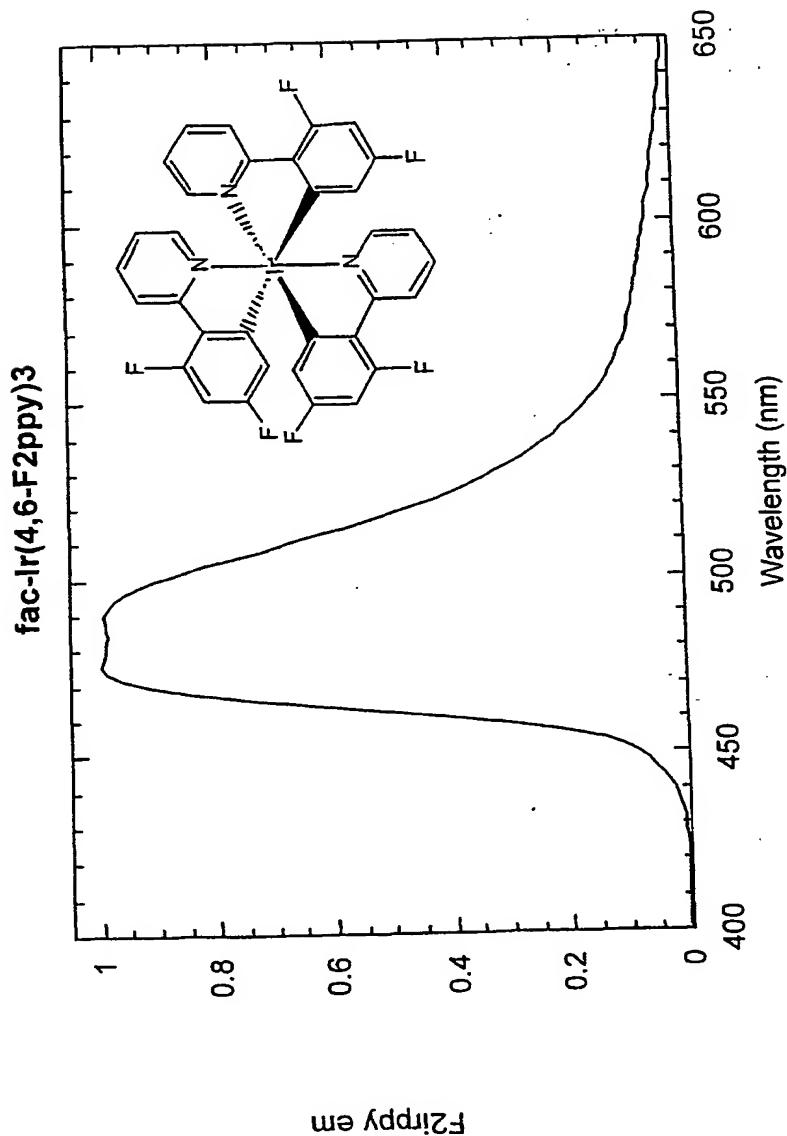


Figure 7q

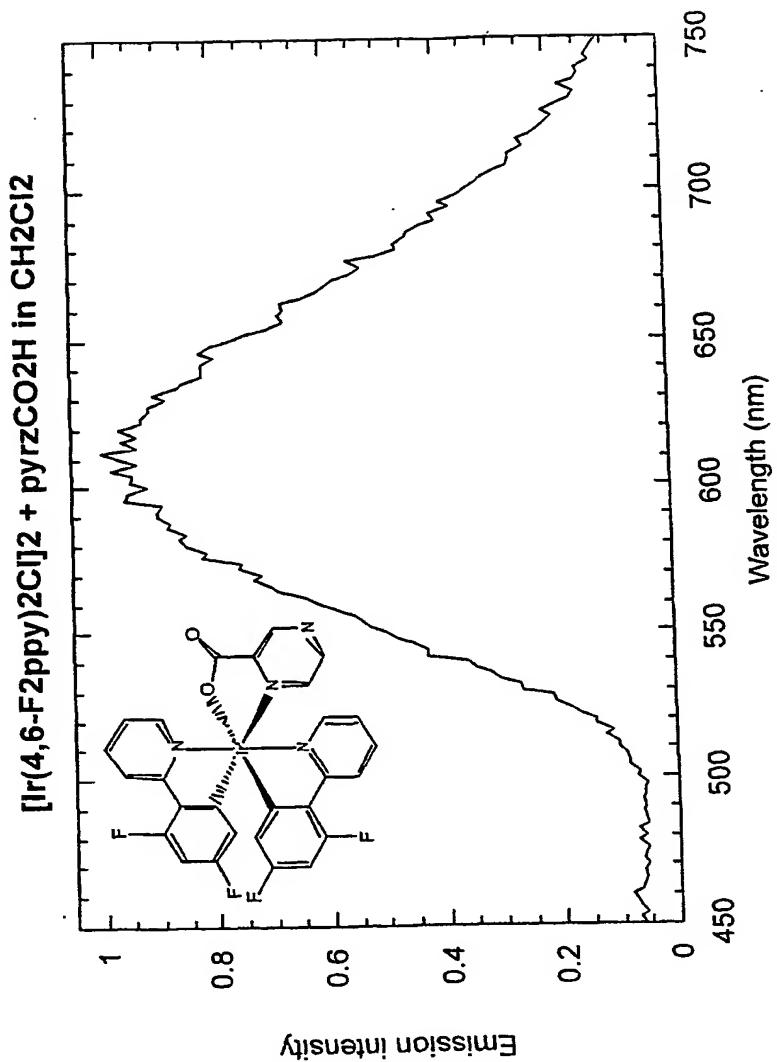


Figure 7r

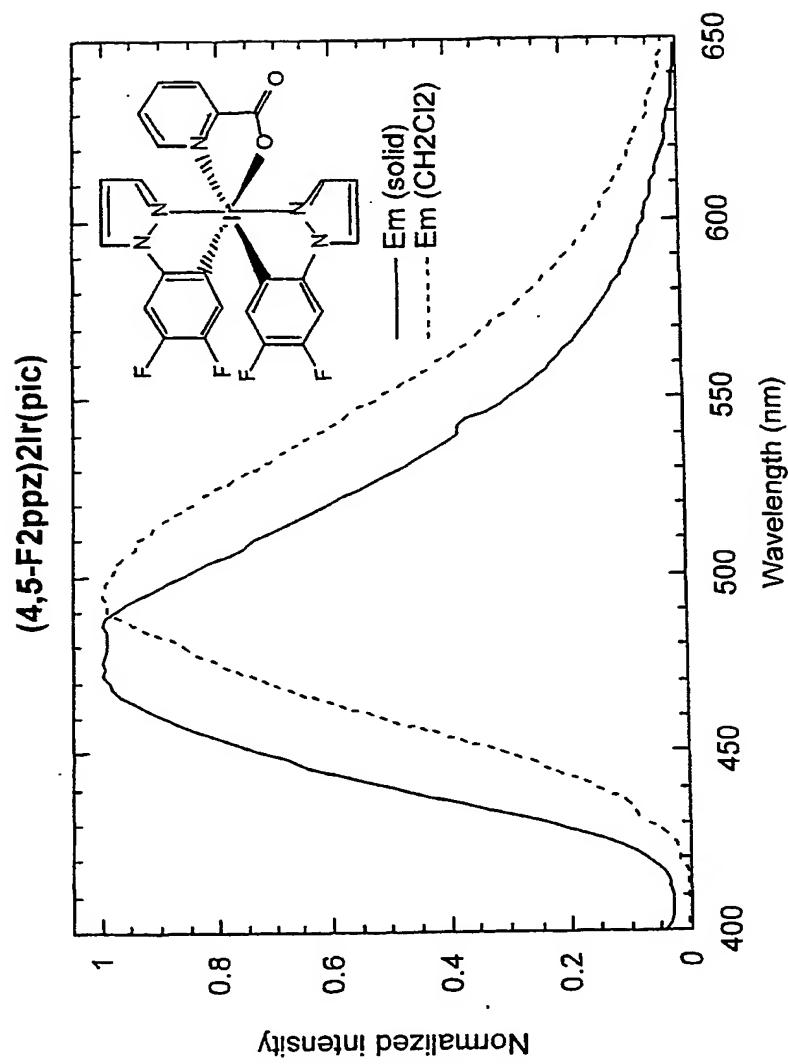


Figure 8a

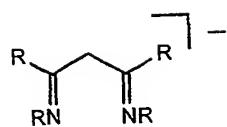
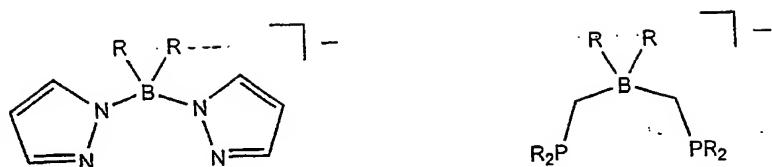
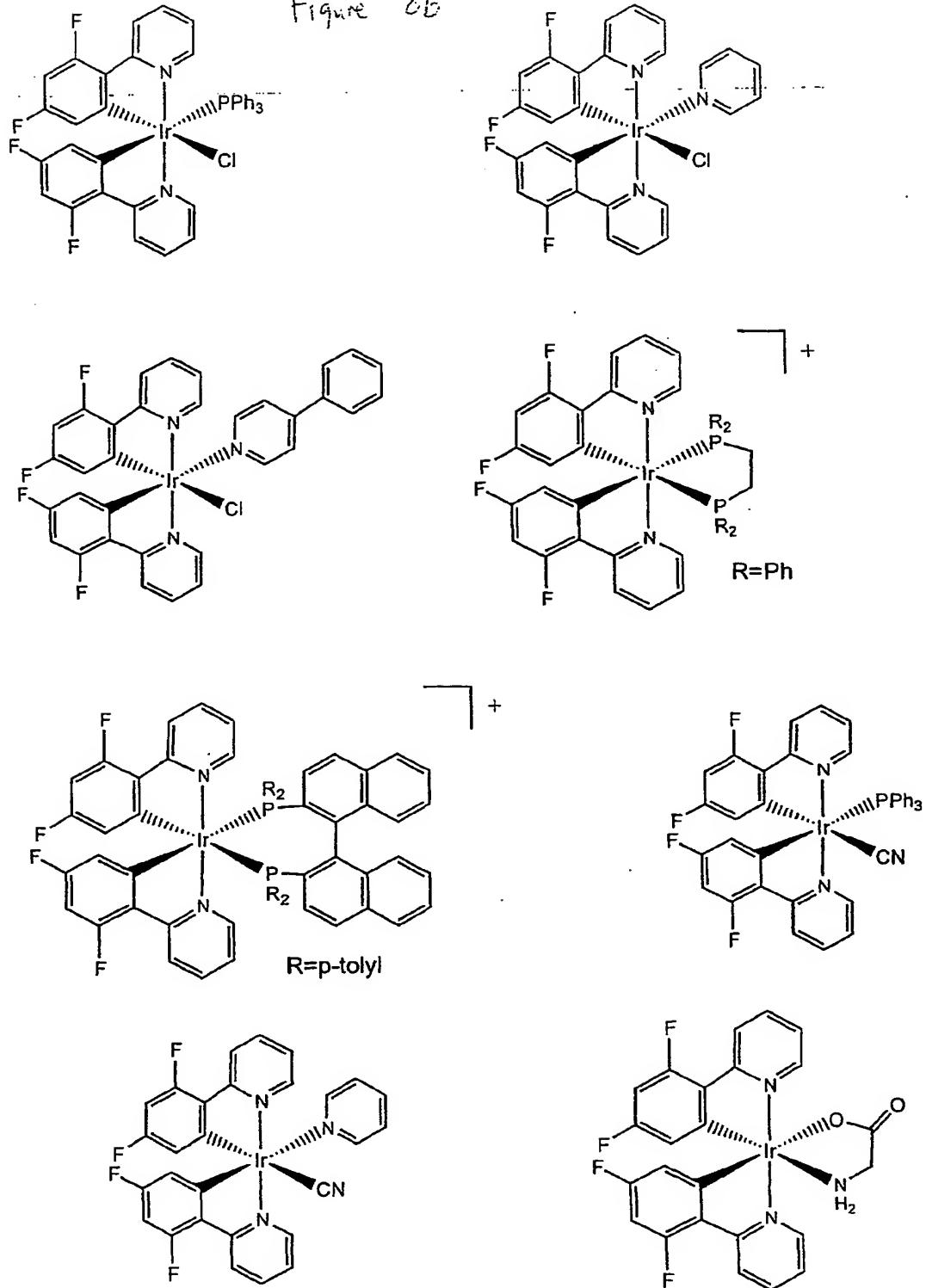


Figure 8b



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Figure

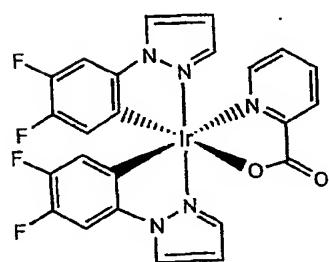
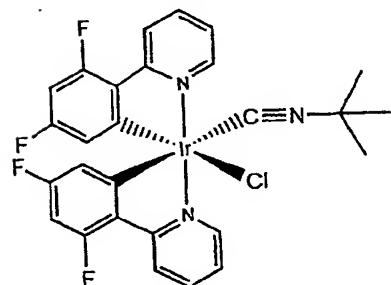
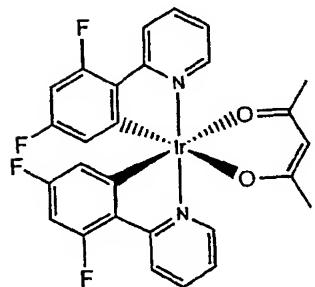
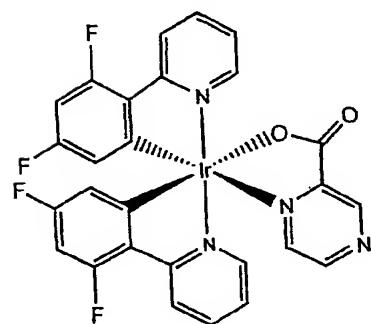
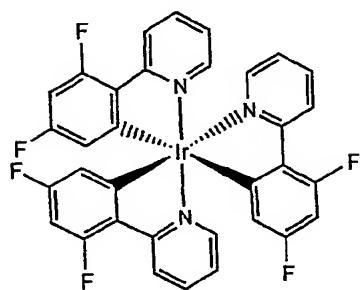
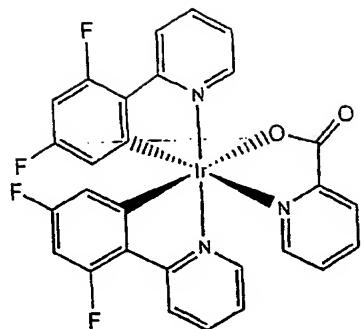
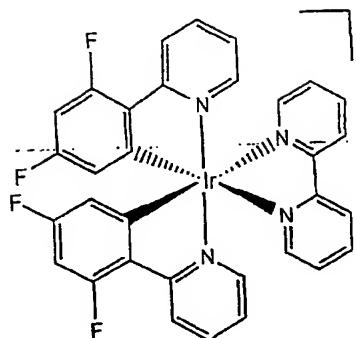
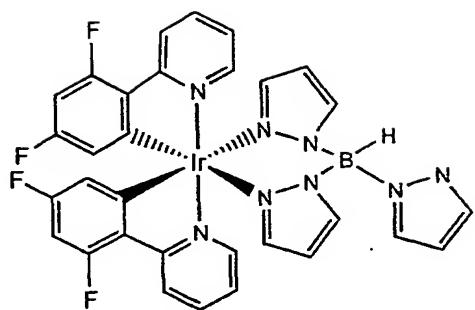
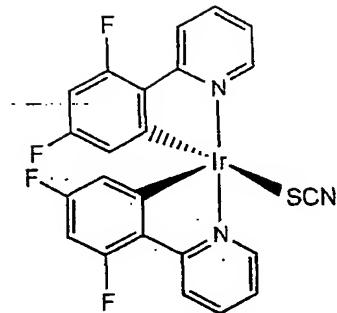
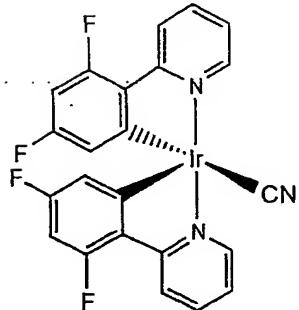
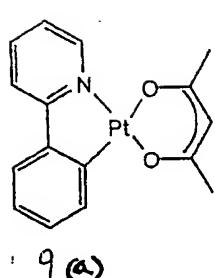


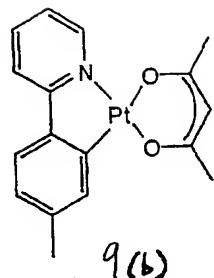
Figure 8d





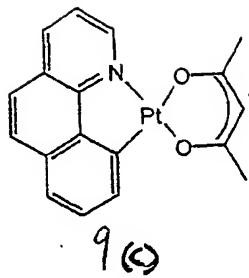
9(a)

(ppy)Pt(acac)



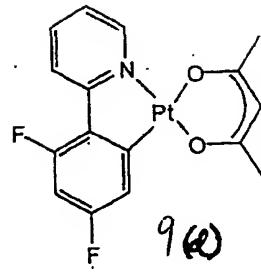
9(b)

(tpy)Pt(acac)

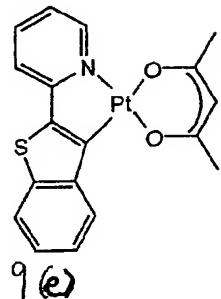


9(c)

(bzq)Pt(acac)

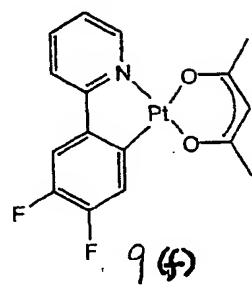


9(d)

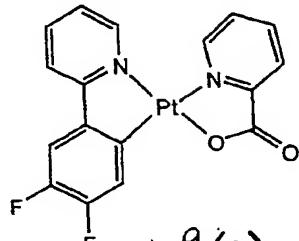
(4,6-F₂ppy)Pt(acac)

9(e)

(btp)Pt(acac)



9(f)

(4,5-F₂ppy)Pt(acac)

9(g)

(4,5-F₂ppy)Pt(pico)

Figures 9(a) - 9(g)

Figure 10: This Emission spectrum shows the spectra of both $\text{Pt}(\text{ppy})_2$ and $\text{Pt}(\text{ppy})_2\text{Br}_2$. The former gives green emission, partly from MLCT transitions, and the latter gives blue emission, predominantly from a triplet $\pi-\pi^*$ transition. The structure observed for the $\text{Pt}(\text{ppy})_2\text{Br}_2$ spectrum is consistent with ligand centered emission. The luminescent lifetimes for the two complexes are 4 and 150 μsec .

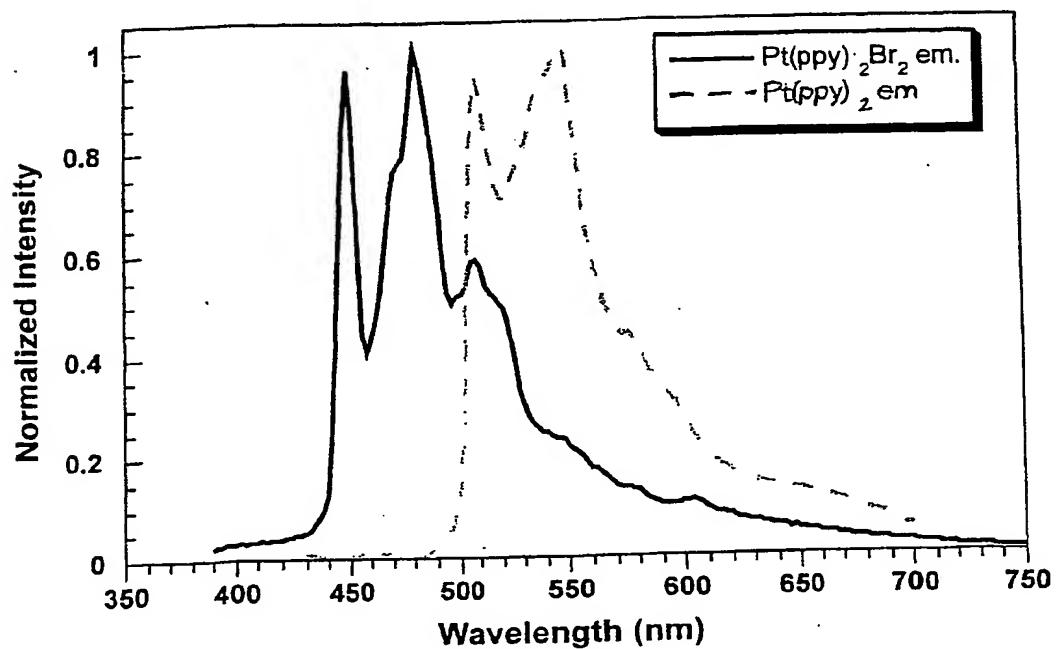


Figure 10

Figure II: This plot shows the emission spectra of $(ppy)AuCl_2$ and $(ppy)Au(2,2'-biphenylene)$. Both emit from ligand triplet $\pi-\pi^*$ transitions.

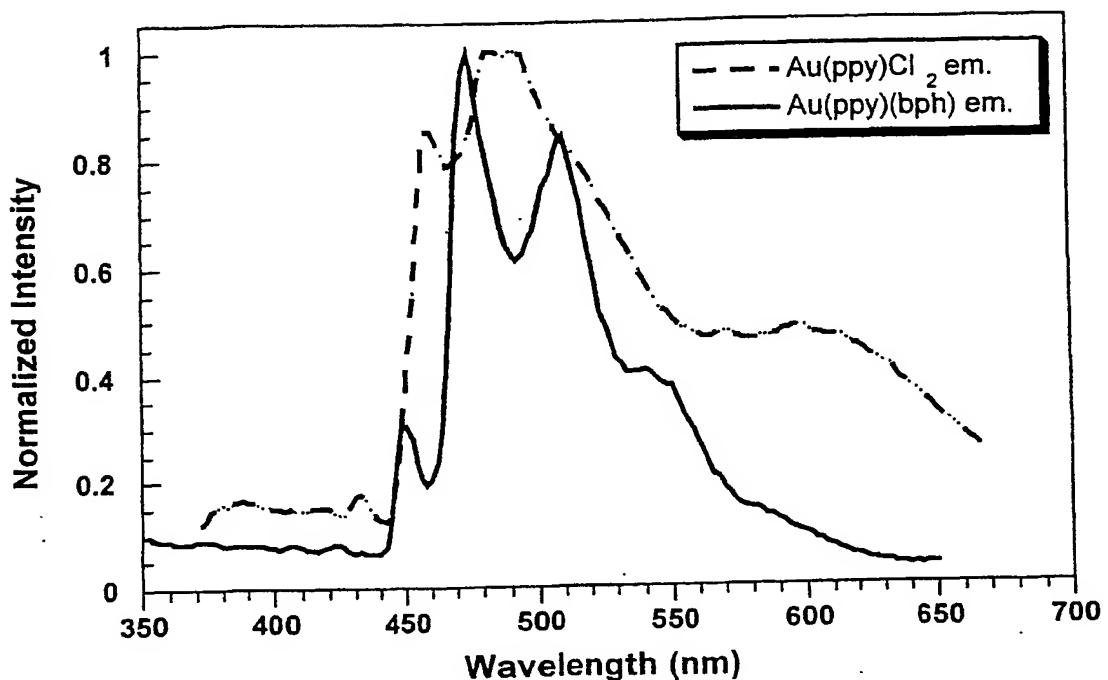


Figure II

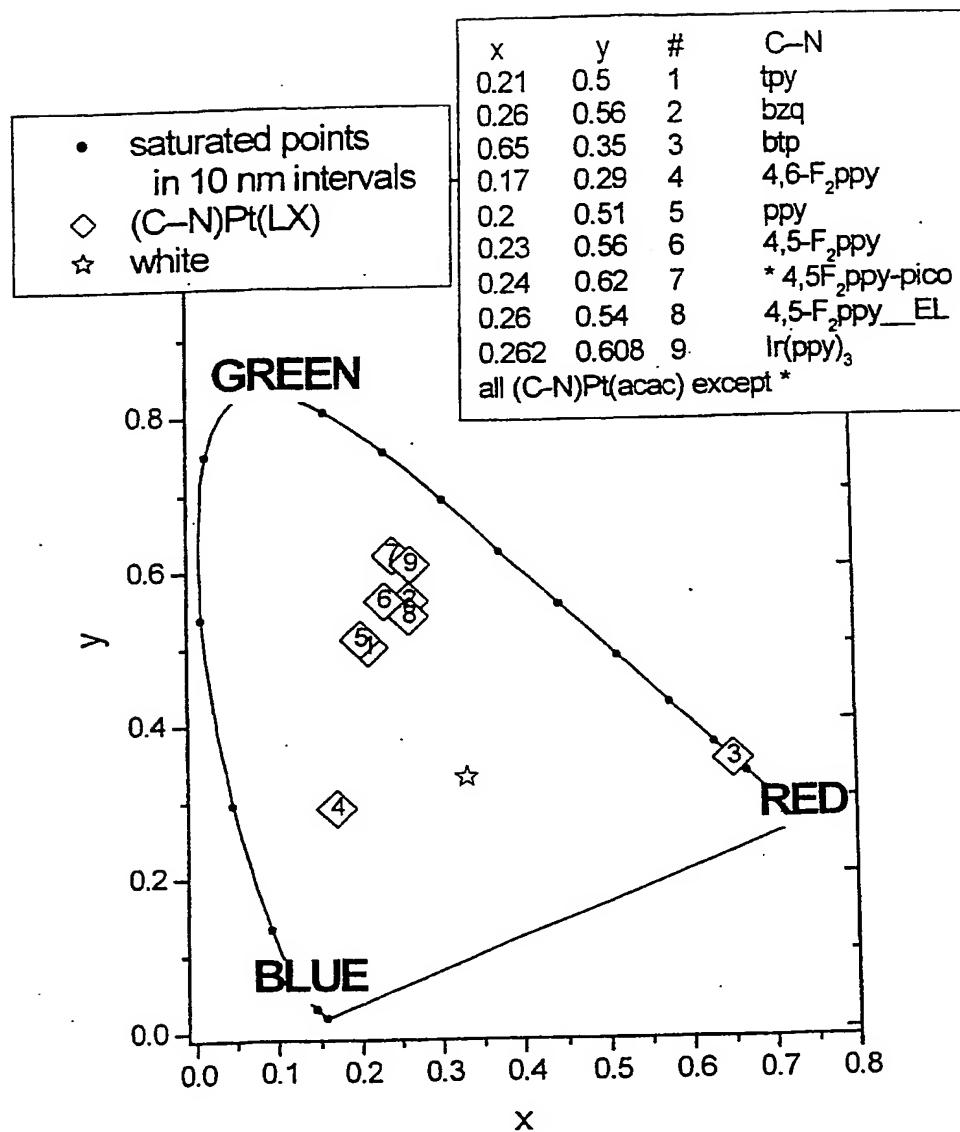


Figure 12

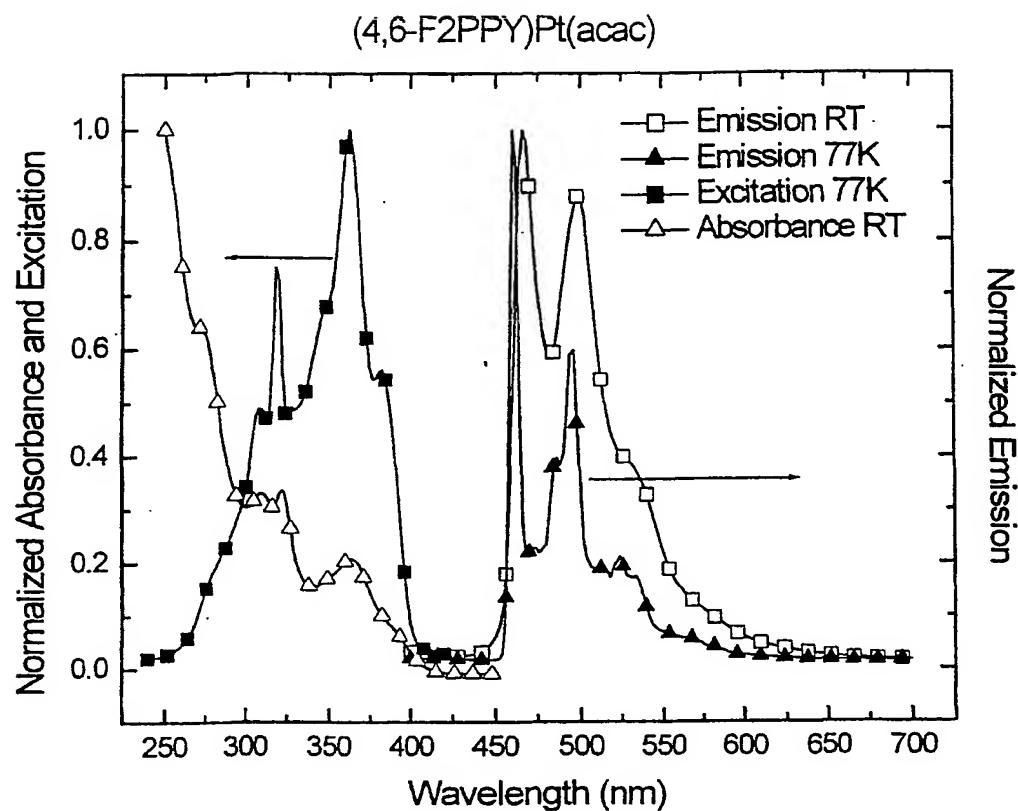


Figure 13

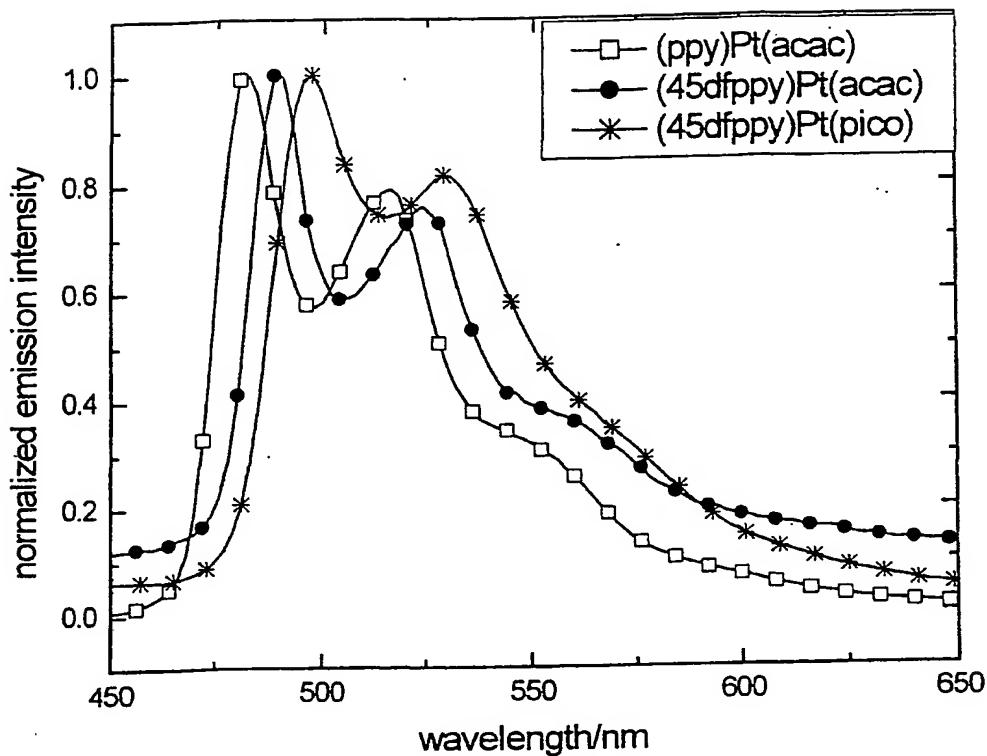


Figure 14

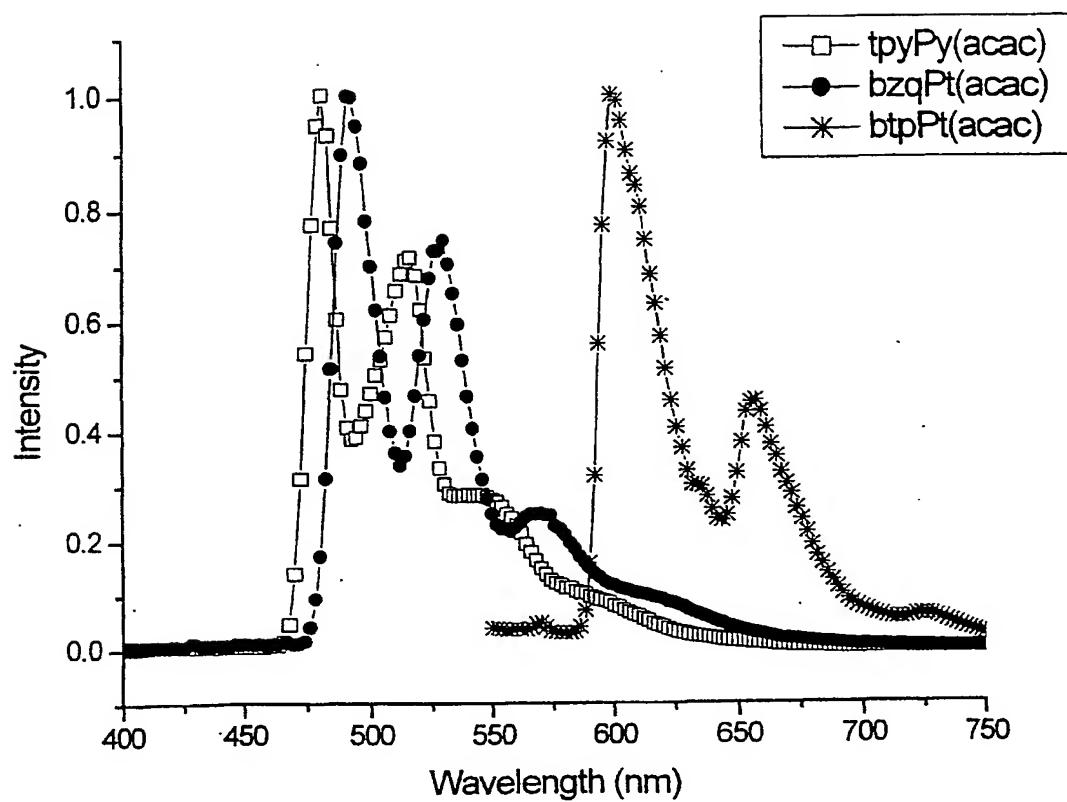


Figure 15

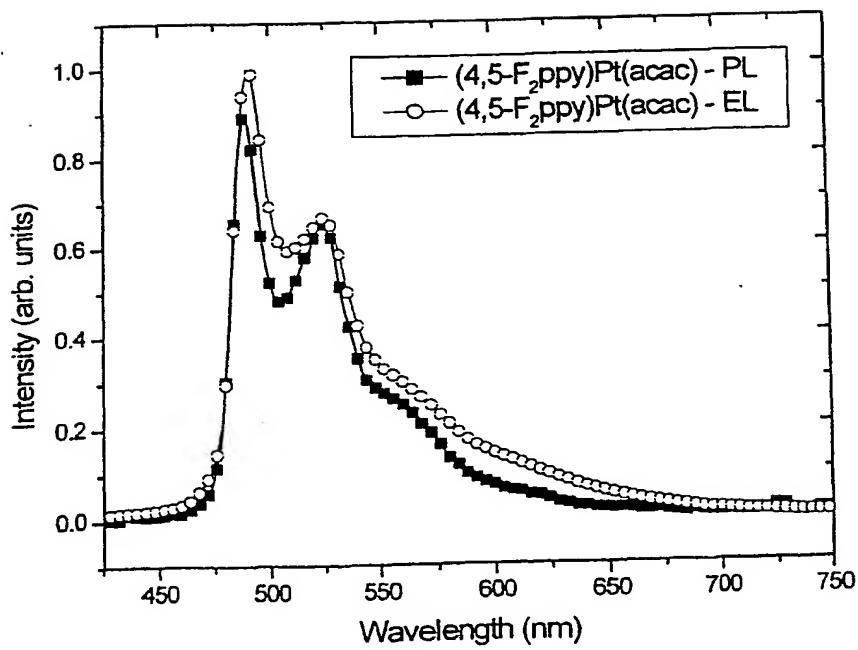


Figure 16

FIG. 17

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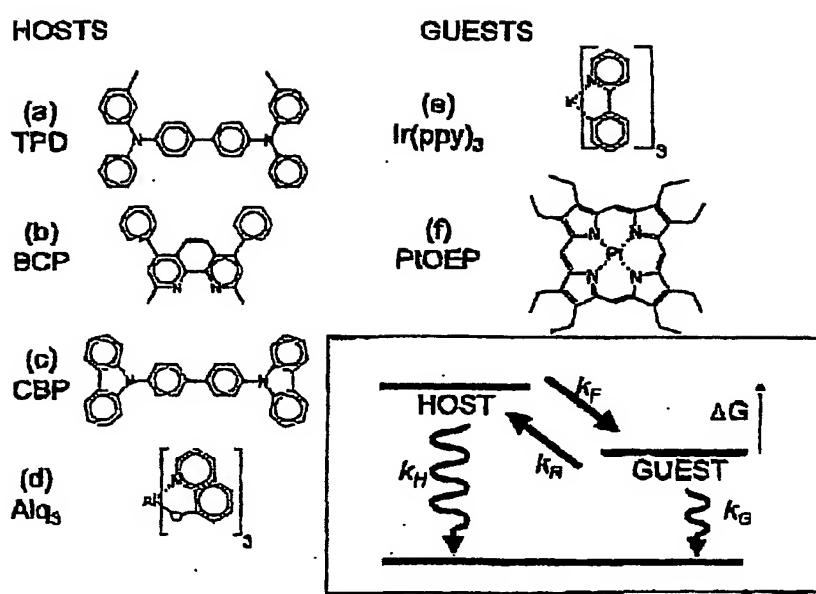


FIG. 18

PRB 62

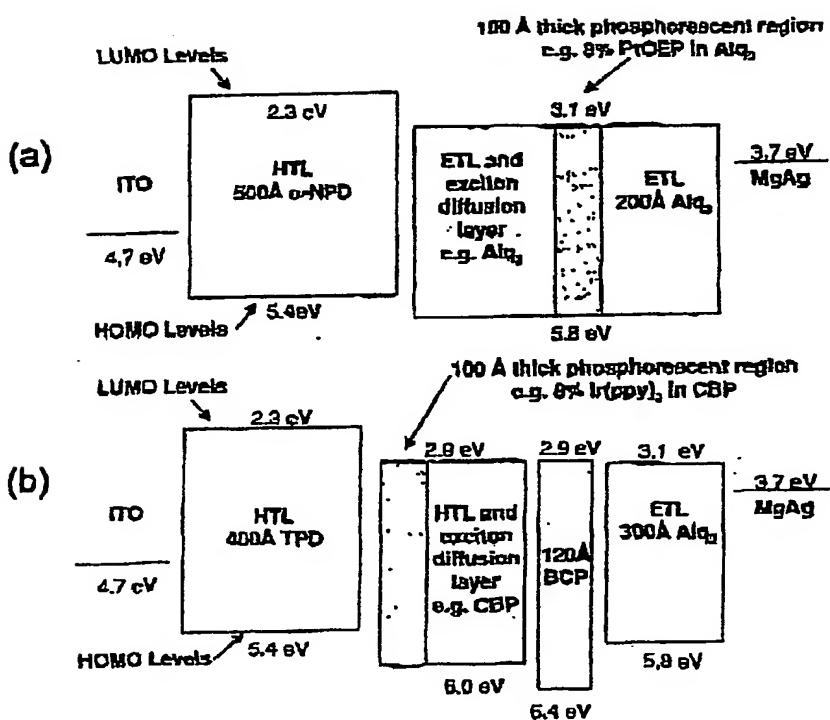


FIG. 19

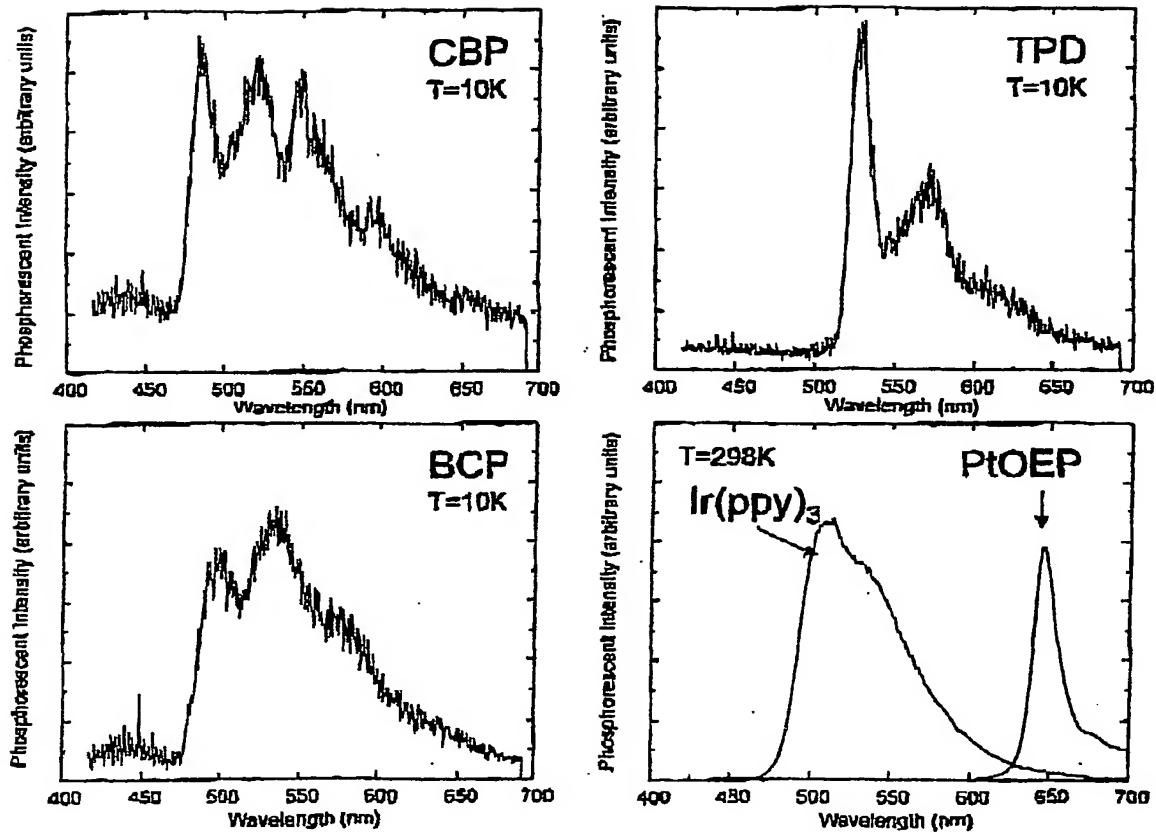


FIG. 20

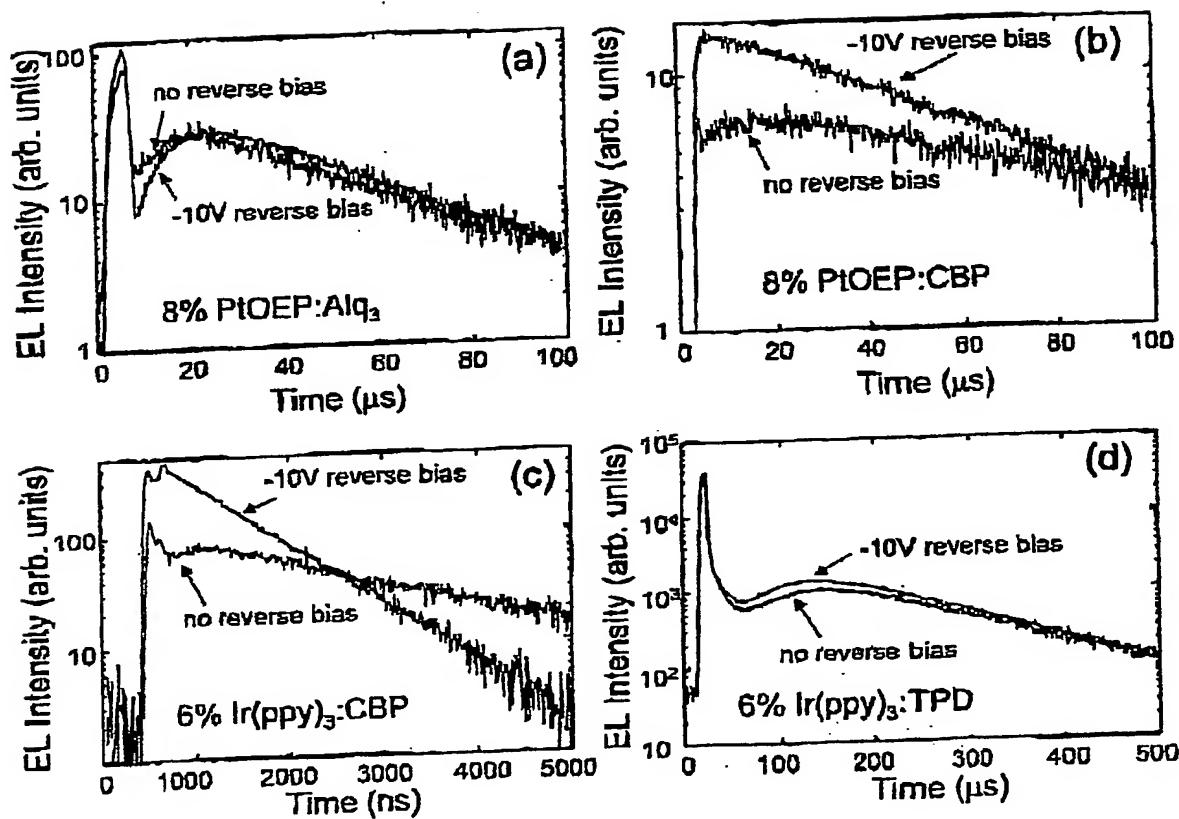


FIG. 21

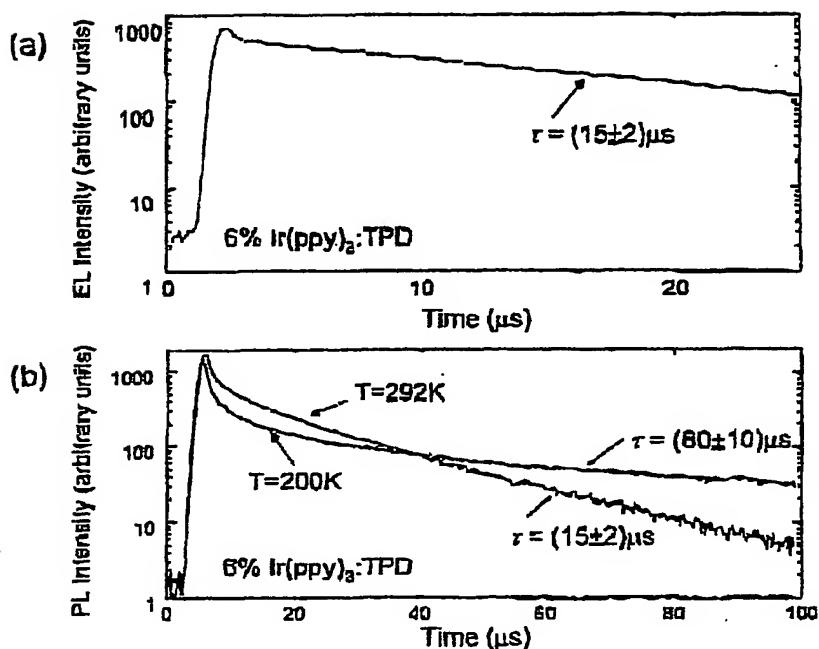


FIG. 22

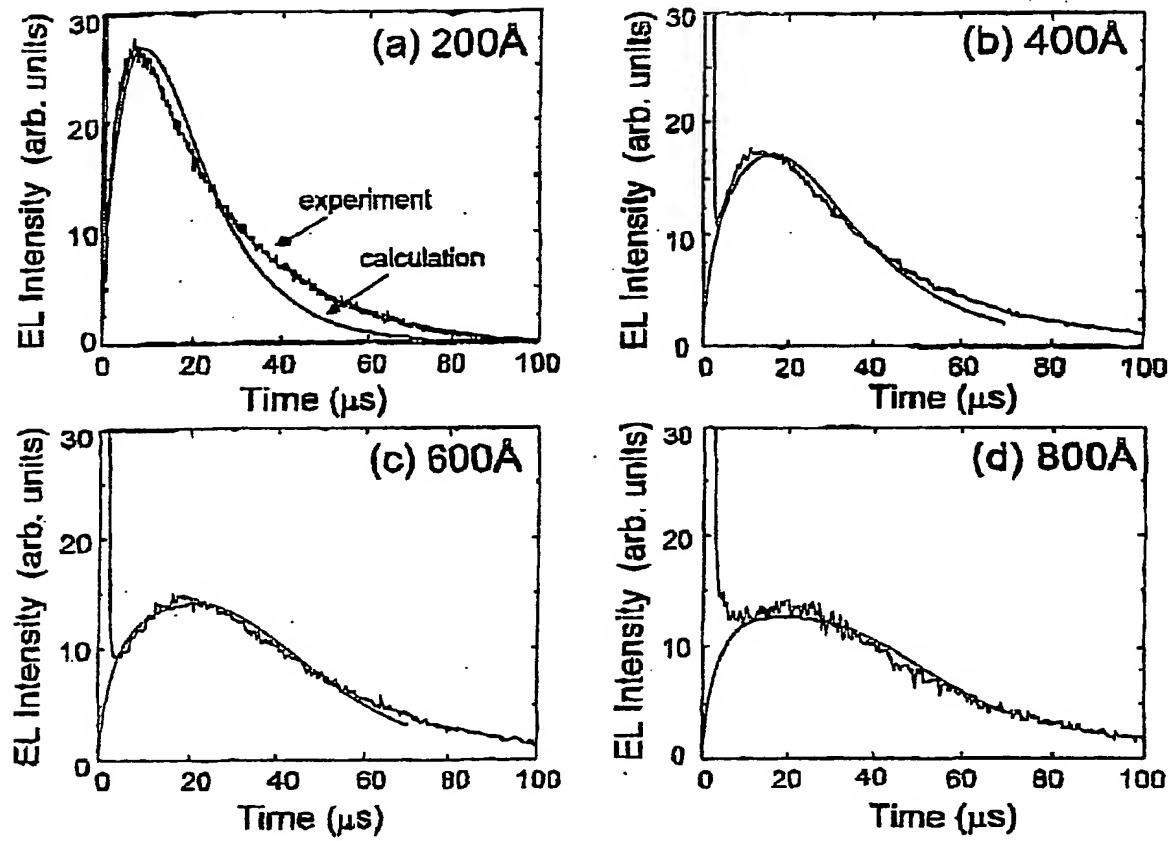


Fig. 23

